



STIC Search Report

Biotech-Chem Library

STIC Database Tracking Number: 10/644021

TO: Chun Crowder
Location: REM-3B59/3C70
Art Unit: 1644
Monday, March 27, 2006

Case Serial Number: 10/644021

From: Deirdre Arnold
Location: Biotech-Chem Library
REM 1A55
Phone: 571-272-2532

Deirdre.Arnold@uspto.gov

Search Notes

Please feel free to contact me if you have any questions or would like to amend the search.

Thank you for using STIC services.

Regards,
Deirdre Arnold

THIS PAGE BLANK (USPTO)

Crowder, Chun

To: STIC-Biotech/ChemLib
Subject: Sequence search for 10/644,021

SEQ ID NO:2 against commercial and interference protein databases.

Thanks!

Chun Crowder, Ph.D.
81042
Patent Examiner
TC1644
Remsen/03B59
3c70
571-272-8142

THIS PAGE BLANK (USPTO)

GenCore version 5.1.7
Copyright (c) 1993 - 2006 Biocceleration Ltd.

Om protein - protein search, using sw model
Run on: March 24, 2006, 16:42:20 ; Search time 187 Seconds
(without alignments)

Scoring table: BL0SUM62
Gapop 10.0 , Gapext 0.5

Title: US-10-644-021a-2
Perfect score: 1952
Sequence: 1 MEFVYKCLGHPEEPYFLNVLRR... .WQYLTTLSQVTEDYVQTGEH 374
878.758 Million cell updates/sec

Searched: 2443163 seqB, 439378781 residues
Total number of hits satisfying chosen parameters: 2443163

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_21;*

1: genesedp1980b;*
2: genesedp2000b;*
3: genesedp2001b;*
4: genesedp2002b;*
5: genesedp2003b;*
6: genesedp2003ab;*
7: genesedp2003bb;*
8: genesedp2004b;*
9: genesedp2005b;*

Pred. No. 1 is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

RESULT 1			
ID	Description	XX	XX
ABG72689;	ABG72689 standard; protein; 374 AA.	AC	AC
		DT	05-MAR-2003 (first entry)
		DE	Human squalene synthase.
		XX	Human; enzyme; squalene synthase; cholesterol-related disease; cardiovascular disease; chromosome 8; SNP; single nucleotide polymorphism.
		KW	
		OS	Homo sapiens.
		PN	US2002142418-A1.
		XX	XX
		PD	03-OCT-2002.
		XX	XX
		PF	29-MAR-2001; 2001US-0082004.
		PR	29-MAR-2001; 2001US-0082004.
		XX	XX
		PA	(WEIM/) WEI M.
		PA	(YANC/) YAN C.
		PA	(DFRA/) DI FRANCESCO V.
		PA	(BEAS/) BEASLEY E M.
		XX	XX
		PT	Wei M, Yan C, Di Francesco V, Beasley EM;
		XX	XX
		DR	WPI; 2003-155945/15.
		DR	N-PSDB; ABX14651, ABX14652.
		XX	XX
		PT	Novel isolated enzyme protein related to synthase enzyme subfamily, useful as models for developing human therapeutic targets, aid in the identification of therapeutic proteins and as immunogens to raise antibodies.
		PT	Claim 1, Fig 2; 76pp; English.
		PS	The invention relates to an isolated enzyme protein (a squalene synthase)

Result No.	Score	Query Match Length	DB ID	Description	ALIGNMENTS
1	1952	100.0	374	6 ABCG2689	ABP72689 Human seqA
2	1952	100.0	374	8 ADNB6662	Actn9662 Novel hum
3	1920.5	98.4	417	2 AAW01739	Aaw01739 Human seqA
4	1920.5	98.4	417	7 ADEB8269	Ades8269 Human Pro
5	1920.5	98.4	417	7 ADD6345	Add6345 Human Pro
6	1920.5	98.4	417	7 ADEB8261	Ades8261 Human Pro
7	1920.5	98.4	417	7 ADEB8265	Ades8265 Human Pro
8	1920.5	98.4	417	9 ADZ0390	Adz0390 Human Pro
9	1920.5	98.4	417	7 ADJ9872	Adj9872 Novel NOV
10	1920.5	98.4	417	8 ADNB9865	Adnb9865 FarnesyL-
11	1920.5	98.4	417	8 ADNB9864	Adnb9864 FarnesyL-
12	1920.5	98.4	417	8 ADT9957	Adt9957 Human seqA
13	1920.5	98.4	417	9 ADY16510	Ady16510 PRO polyp
14	1920.5	98.4	417	9 ADT0390	Adt0390 Human pro
15	1915.5	98.1	417	2 AAR52605	Aar52605 Human seqA
16	1915.5	98.1	417	8 ADNB9866	Adnb9866 FarnesyL-
17	1915.5	98.1	417	8 ADT9956	Adt9956 FarnesyL-
18	1912.5	98.0	417	8 AD05028	Ad05028 Human seqA
19	1737	89.0	416	5 ABB57061	Abb57061 Mouse isoC
20	1692	86.7	416	7 ADB83186	Adb83186 Rat squal
21	1692	86.7	416	7 ADB8271	Adb8271 Rat prote
22	1692	86.7	416	7 ADEB8259	Adeb8259 Rat Prote
23	1692	86.7	416	7 ADB58267	Adb58267 Rat Prote
24	1692	86.7	416	7 ADE58263	Ades58263 Rat Prote

CC related to synthase enzyme subfamily appearing as ABG72689, its allelic under variant or orthologue and encoded by a nucleic acid that hybridizes under stringency to opposite strand of the cDNA and gene appearing as ABX14651 and ABX14652, or its fragment of a sequence with 70% sequence similarity to the protein. Also disclosed are nucleic acids encoding the protein, antibodies, expression vectors, transformed host cells, transgenic animals, gene chips and identifying modulators/binding agents of the protein or nucleic acid. A pharmaceutical composition comprising the protein is useful for treating a disease or condition mediated by e.g. cholesterol-related diseases and cardiovascular diseases. The protein is useful to identify therapeutics, binding partners and modulating agents. The antibody is useful for identifying, isolating and purifying the protein. The nucleic acids is useful as probes, primers, and chemical intermediates in biological assay. The gene for the synthase enzyme is located on chromosome 8. The present sequence represents the human squalene synthase

SQ Sequence 374 AA;

Query Match	100.0%	Score 1952; DB 6; Length 374;
Best Local Similarity	100.0%	Pred. No. 2, 6e-15;
Matches	374;	Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	1	MEFVKCLGHPEEPFVNVLRRIGKRKVVKMRKDODSLSLTKCYKLQNTSRSPAVTQA 60
Db	1	MEFKVCLGHPEEPFVNVLRRIGKRKVVKMRKDODSLSLTKCYKLQNTSRSPAVTQA 60
Db	1	MEFKVCLGHPEEPFVNVLRRIGKRKVVKMRKDODSLSLTKCYKLQNTSRSPAVTQA 60
QY	61	LDGMRNACVCFYVTLRAIDLDEDMTISVEKKVPLLNHFSFLYQPDWRFMESKEKORQ 120
Db	61	LDGMRNACVCFYVTLRAIDLDEDMTISVEKKVPLLNHFSFLYQPDWRFMESKEKORQ 120
QY	121	VLEDFTPTCHYVAGLVGLSLRSLSSAEEDPVLGEDTERANGLFLQKTNIRDYED 180
Db	121	VLEDFTPTCHYVAGLVGLSLRSLSSAEEDPVLGEDTERANGLFLQKTNIRDYED 180
QY	181	QQSGREFWPOEVNSRYVKVKGDFAKPENIDLAVOCLNLITNALHHIPDVITVLSRLNQ 240
Db	181	QQSGREFWPOEVNSRYVKVKGDFAKPENIDLAVOCLNLITNALHHIPDVITVLSRLNQ 240
QY	241	SVFNCFAIPOVMATLAACYNNOQVFKCAVKRGQAVTLMDDATNNPAKAVIYQME 300
Db	241	SVFNCFAIPOVMATLAACYNNOQVFKCAVKRGQAVTLMDDATNNPAKAVIYQME 300
QY	301	EIHTRIPSDPSSKTRQISTRTQNLNCOLISRSWSPIVLFVMLAALSWQYLT 360
Db	301	EIHTRIPSDPSSKTRQISTRTQNLNCOLISRSWSPIVLFVMLAALSWQYLT 360
QY	361	LSQVTEVDYVQTGEH 374
Db	361	LSQVTEVDYVQTGEH 374

RESULT 2

ADN96862 standard; protein; 374 AA.

ID ADN96862;

AC ADN96862;

DT 26-AUG-2004 (first entry)

DE Novel human enzyme.

XX disease diagnosis; gene expression associated disorder; gene expression; KW disease diagnosis; gene expression associated disorder; gene expression; enzyme peptide; human; enzyme.

OS Homo sapiens.

XX

Key Location/Qualifiers

FT Domain .25

FT /note = Amidation site

FT Domain .41

FT /note = Protein kinase C phosphorylation site

FT Domain .51

Sequence 374 AA;

FT Domain /note = N-glycosylation site

FT Domain /note = Protein kinase C phosphorylation site

FT Domain /note = Casein kinase II phosphorylation site

FT Domain /note = Casein kinase II phosphorylation site

FT Domain /note = Squalene and phyoene synthases signature 1

FT Domain /note = N-myristoylation site

FT Domain /note = Casein kinase II phosphorylation site

FT Domain /note = Casein kinase II phosphorylation site

FT Domain /note = Squalene and phyoene synthases signature 2

FT Domain /note = N-glycosylation site

FT Domain /note = N-myristoylation site

FT Domain /note = Protein kinase C phosphorylation site

FT Domain /note = Protein kinase C phosphorylation site

FT Domain /note = Protein kinase C phosphorylation site

FT Domain /note = Protein kinase C phosphorylation site

FT Domain /note = Protein kinase C phosphorylation site

FT Domain /note = Protein kinase C phosphorylation site

FT Domain /note = Protein kinase C phosphorylation site

PR 29-MAR-2001; 2001US-00820004.

PA (APPL-) APPLEA CORP.

PI Wei M, yan C, Di Francesco V, Beasley E;

XX DR WPT; 2004-419461/39.

XX DR N-PSDB; ADN96861, ADN96863.

PT New isolated enzyme proteins, useful for diagnosing or treating diseases characterized by absence, inappropriate, or unwanted expression of the protein, or as a reagent in assays for determining levels of protein in biological sample.

XX PT

PS Claim 1; SEQ ID NO 2; 76pp; English.

XX

CC The invention describes an isoLATED enzyme peptide (I) comprising a defined sequence of 374 amino acids. Also described are: an isolated antibody that selectively binds to (I); a method for producing the peptides; a method for detecting the presence of the peptides; a method for identifying a modulator of the peptide; a method for identifying an agent that binds to the peptide; a pharmaceutical composition comprising an agent identified by the method of (5) and a pharmaceutical carrier; a method for treating a disease or condition mediated by a human enzyme; a method for identifying a modulator of the expression of the peptide; and an isolated human enzyme peptide having an amino acid sequence that shares at least '70% homology with SEQ ID NO. 2. Specifically claimed is an enzyme peptide comprising 374 amino acids (SEQ ID NO. 2). The peptides are useful for substantial or specific assays, e.g. biological, or drug screening assays, as a reagent in assay for determining levels of protein in biological sample; and as markers for tissues where the corresponding protein is expressed. It can also be used to screen a compound for the ability to stimulate or inhibit interaction between the enzyme protein and a molecule that normally interacts with the enzyme protein. They are also useful as a target for diagnosing a disease or predisposition to disease mediated by the peptide. It can also be used for treating disorders characterised by absence, inappropriate, or unwanted expression of the protein. This is the amino acid sequence of the novel human enzyme peptide of the invention.

CC Sequence 374 AA;

Query Match 100.0%; Score 1952; DB 8; Length 374;
 Best Local Similarity 100.0%; Pred. No. 2.6e-195; Mismatches 0; Indels 0; Gaps 0;
 Matches 374; Conservative 0; PDB ID 0; SRS 0;

QY 1 MERVKCLGHPESFYNLVRIGRKPKMPKMDPSLSSLKTCYKVNQTSRSFAVIOA 60
 1 MERVKCLGHPESFYNLVRIGRKPKMPKMDPSLSSLKTCYKVNQTSRSFAVIOA 60
 Db 61 LDGEMRNACVIFTYLVRALDTLDDMTISVEKKVPLHNPHSTLYQDWRMFKESKEKDRQ 120
 QY 61 LDGEMRNACVIFTYLVRALDTLDDMTISVEKKVPLHNPHSTLYQDWRMFKESKEKDRQ 120
 61 LDGEMRNACVIFTYLVRALDTLDDMTISVEKKVPLHNPHSTLYQDWRMFKESKEKDRQ 120
 Db 121 VLEDPTTYCHYVAGLVLGQLSRLFSASEFDPLVGDFTERANSGLFLQKTNIRDYLED 180
 QY 121 VLEDPTTYCHYVAGLVLGQLSRLFSASEFDPLVGDFTERANSGLFLQKTNIRDYLED 180
 Db 121 VLEDPTTYCHYVAGLVLGQLSRLFSASEFDPLVGDFTERANSGLFLQKTNIRDYLED 180
 QY 181 QSGREFWQBVNSVRYKLGDAKPNIDLAVQCLNELITNALHHFDPVITLRSRNO 240
 Db 181 QSGREFWQBVNSVRYKLGDAKPNIDLAVQCLNELITNALHHFDPVITLRSRNO 240
 QY 241 SVFNFCATPQVMAITALAACYNNOQVFKGAVKRGKQAVTLMMDATNNPAVKAIYQME 300
 Db 241 SVFNFCATPQVMAITALAACYNNOQVFKGAVKRGKQAVTLMMDATNNPAVKAIYQME 300
 301 EYMRIPSDPSKSSTKROISTRTQNLPNCQLISRSRSHSPIVLSFVMLLAALSQWQYLT 360
 Db 301 EYMRIPSDPSKSSTKROISTRTQNLPNCQLISRSRSHSPIVLSFVMLLAALSQWQYLT 360
 QY 361 LSQTEDYVQTGEH 374
 Db 361 LSQTEDYVQTGEH 374

RESULT 3
 AAW01739
 ID AAW01739 standard; protein; 417 AA.
 XX
 AC AAW01739;
 DT 25-MAR-2003 (revised)
 DT 17-APR-1997 (first entry)
 XX Human squalene synthetase.
 XX
 KW squalene synthetase; Saccharomyces cerevisiae; Erg9; biosynthesis; recombinant production; soluble; sterol.
 KW farneyl pyrophosphate; mevalonate production; characterisation; study; recombinant production; soluble; sterol.
 OS Homo sapiens.
 PN USS589372-A
 XX 31-DEC-1996.
 PD 08-DEC-1994; 94US-00351981.
 PR 26-SEP-1990; 90US-00588235.
 PR 10-JUL-1992; 92US-00911835.
 PA (SQUI) SQUIBB & SONS INC E R.
 PI Robinson GW;
 DR WPI; 1997-076848/07.
 DR N-PSDB; AAT59298.

102(6)

the first committed enzyme of sterol biosynthesis, has a low affinity for farneyl pyrophosphate (FPP) which is incorporated into a variety of end-products e.g. dolichols, ubiquinone, hormones, haem A, sterols and some isoprenylated proteins. Differential synthesis of the FPP-derived products is controlled through both regulated enzyme synthesis and differing affinities for FPP. Squalene synthetase levels are regulated, a ten-fold depression in activity is seen in cells when the accumulate sufficient cholesterol. Together, these factors, in concert with regulation of mevalonate production by HMG CoA reductase (HMGCR), ensure that adequate non-sterol product of FPP are made both in cells actively synthesising cholesterol from FPP and in cells that receive most of their cholesterol exogenously, through uptake mediated by the low density lipoprotein receptor. The characterisation of squalene synthetase and its recombinant production (esp. in solubilised form) would be useful to expedite its study. (Updated on 25-MAR-2003 to correct PF field.)

CC SQ Sequence 417 AA;

Query Match 98.4%; Score 1920.5; DB 2; Length 417;
 Best Local Similarity 89.7%; Pred. No. 6.1e-192; Mismatches 0; Indels 43; Gaps 1; Matches 374; Conservative 0; PDB ID 0; SRS 0;

QY 1 MERVKCLGHPESFYNLVRIGRKPKMPKMDPSLSSLKTCYKVNQTSRSFAVIOA 60
 1 MERVKCLGHPESFYNLVRIGRKPKMPKMDPSLSSLKTCYKVNQTSRSFAVIOA 60
 Db 61 LDGEMRNACVIFTYLVRALDTLDDMTISVEKKVPLHNPHSTLYQDWRMFKESKEKDRQ 120
 QY 61 LDGEMRNACVIFTYLVRALDTLDDMTISVEKKVPLHNPHSTLYQDWRMFKESKEKDRQ 120
 61 LDGEMRNACVIFTYLVRALDTLDDMTISVEKKVPLHNPHSTLYQDWRMFKESKEKDRQ 120
 Db 121 VLEDPTTYCHYVAGLVLGQLSRLFSASEFDPLVGDFTERANSGLFLQKTNIRDYLED -CHYVAGL 137
 121 VLEDPTTYCHYVAGLVLGQLSRLFSASEFDPLVGDFTERANSGLFLQKTNIRDYLED -CHYVAGL 180
 QY 138 IGSLRFLFSASEFDPLVGDFTERANSGLFLQKTNIRDYLEDQSGREFWQBVNSVRY 197
 Db 181 IGSLRFLFSASEFDPLVGDFTERANSGLFLQKTNIRDYLEDQSGREFWQBVNSVRY 240
 QY 198 KKLGDFAKEPENIDLAVQCLNELITNALHHFDPVITLRSRNOVSFNFCATPQVMAITAL 257
 Db 241 KKLGDFAKEPENIDLAVQCLNELITNALHHFDPVITLRSRNOVSFNFCATPQVMAITAL 300
 258 AACTNQQVFKGAVKRGKQAVTLMMDATNNPAVKAIYQMEIYHRIPSDPSKSCTR 317
 Db 301 AACTNQQVFKGAVKRGKQAVTLMMDATNNPAVKAIYQMEIYHRIPSDPSKSCTR 360
 QY 318 QIISTIRTQNLPNCQLISRSRSHSPIVLSFVMLLAALSQWQYLTLSQVTDYVQTGEH 374
 Db 361 QIISTIRTQNLPNCQLISRSRSHSPIVLSFVMLLAALSQWQYLTLSQVTDYVQTGEH 417

RESULT 4
 AD58269
 ID AD58269 standard; protein; 417 AA.
 XX
 AC AD58269;
 XX
 DT 29-JAN-2004 (first entry)
 XX Human Protein P37268, SEQ ID NO 4140.
 DB Human; pain; neuronal tissue; gene therapy; Human; pain; neuronal tissue; gene therapy; chronic constriction injury; cci; Human; pain; neuronal tissue; gene therapy; chronic constriction injury; cci; Human; pain; neuronal tissue; gene therapy; spared nerve injury; sni; Chung.
 XX Homo sapiens.
 XX WO2003016475-A2.
 PT 27-FEB-2003.
 XX
 PS Claim 1; Fig 9; 42pp; English.
 XX
 CC This sequence shows the Human squalene synthetase. Squalene synthetase,

PR	14-AUG-2001; 2001US-0312147P.	360	301 AACINNQQVFGAVAKIRKG@AVTLMADATNNPAKAIYQWEEIYHRIPPSDPSSKTR
PR	01-NOV-2001; 2001US-0346382P.	361	318 QIISTIRTQNPNCOLISRSYHSPYLFSVMLAALSWQYTTLTSQTYEDVOTGEH
PR	26-NOV-2001; 2001US-033347P.	374	
XX		361	QIISTIRTQNPNCOLISRSYHSPYLFSVMLAALSWQYTTLTSQTYEDVOTGEH 417
PA	(GEHO) GEN HOSPITAL CORP.		
PA	(FARB) BAYER AG.		
XX			
PT	Woolf C., D'urso D., Befort K., Costigan M,		
PT	WPI; 2003-268312/26.		
PS	GENBANK; P37268.		
PT	New composition comprising two or more isolated polypeptides, useful for preparing a medicament for treating pain in an animal.		
PT	Claim 1; Page; 1017pp; English.		
XX			
CC	The invention discloses a composition comprising two or more isolated rat or human polynucleotides or a polynucleotide which represents a fragment, derivative or allelic variation of the nucleic acid sequence. Also claimed are a vector comprising the novel polynucleotide, a host cell comprising the vector, a method for identifying a nucleotide sequence which is differentially regulated in an animal subjected to pain and a kit to perform the method, an array, a method for identifying an agent that increases or decreases the expression of the polynucleotide sequence that is differentially expressed in neuronal tissue of a first animal subjected to pain, a method for identifying a compound which regulates the expression of a polynucleotide sequence which is differentially expressed in an animal subjected to pain, a method for identifying a compound that regulates the activity of one or more of the polynucleotides, a method for producing a pharmaceutical composition, a method for identifying a compound or small molecule that regulates the activity in an animal of one or more of the polypeptides given in the specification, a method for identifying a compound useful in treating pain and a pharmaceutical composition comprising the one or more polypeptides or their antibodies. The polynucleotide or the compound modulates its activity is useful for preparing a medicament for treating pain (e.g. spinal segmental nerve injury (Chung), chronic constriction injury (CCI) and spared nerve injury (SNI)) in an animal (e.g. gene therapy). The sequence presented is a human protein (shown in Table 2 of the specification) which is differentially expressed during pain. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic form directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.		
XX			
SQ	Sequence 417 AA:		
Query Match	98.4%; Score 1920.5; DB 7; Length 417;		
Best Local Similarity	89.7%; Pred. No. 6.1e-192;		
Matches	374; Conservative 0; Mismatches 0; Indels 43; Gaps 1;		
QY	1 MERVKCLGHPEEFNLVRIGRKVMMKDQDLSLSSIAKTCYKLNGTSRSFAVIOA 60		
DB	1 MERVVKCLGHPEEFNLVRIGRKVMMKDQDLSLSSIAKTCYKLNGTSRSFAVIOA 60		
QY	61 LDECBMRNAVCIFYVTLRDLTDLDMTISVEKKVPLLRPHSFYQPDFRMESKEKDRQ 120		
DB	61 LDGEMRMRNAVCIFYVTLRDLTDLDMTISVEKKVPLLRPHSFYQPDFRMESKEKDRQ 120		
QY	121 VLEDPPF-----YCHTAGLV 137		
DB	121 VLEDPPF-----YCHTAGLV 180		
QY	138 IGSRLFSASEFEDPLVGDETERANSMGLQKNTIRYLEDQOGGRFSEFWWSRV 197		
DB	181 IGSLRFSASEFEDPLVGDETERANSMGLQKNTIRYLEDQOGGRFSEFWWSRV 240		
QY	198 KKLGDFAKPEPNIDLAQCNELITNAHHPDVITYLSLURNOVENFCAPQMAIAT 257		
DB	241 KKLGDFAKPEPNIDLAQCNELITNAHHPDVITYLSLURNOVNFCAPQMAIAT 300		
QY	258 AACYNQNQVFKGAVKIRKGQAVTMDATNNPAKAIYQWEEIYHRIPPSDPSSKTR 317		

XX SQ Sequence 417 AA;

XX Query Match 98.4%; Score 1920.5; DB 7; Length 417; Best Local Similarity 89.7%; Pred. No. 6_1e-192; Matches 374; Conservative 0; Mismatches 0; Indels 43; Gaps 1;

OY 1 MEFVKCLGHPEEFYNUFRIGRKPKMVKMPQDSSLSLKTCYKYNQTSRSFAAVIA 60
Db 1 MEFVKCLGHPEEFYNUFRIGRKPKMVKMPQDSSLSLKTCYKYNQTSRSFAAVIA 60OY 61 LDGEMRNACIVCFIYLVRALDTEDDMTISVEKKVPLJNHFHFLYQDWRFNEKDRQ 120
Db 61 LDGEMRNACIVCFIYLVRALDTEDDMTISVEKKVPLJNHFHFLYQDWRFNEKDRQ 120OY 121 VLEDPTP----- -YCHYAGLV 137
Db 121 VLEDPTPISLEFRNLAEKYQTADICRMGIGMAEFLDKHVTSEQEDWKCHYAGLV 180OY 138 IGSLRLFSAEFDPLVGDTERANSMGFLQTKNTIRDYLEDQGGERFWQEWRSRYV 197
Db 181 IGSLRLFSAEFDPLVGDTERANSMGFLQTKNTIRDYLEDQGGERFWQEWRSRYV 240OY 198 KKLQGDFAKERENIDLAQOCNELTNAHHIPDVITYSRLRQSVFNCAIQPQWMAATL 257
Db 241 KKLQGDFAKERENIDLAQOCNELTNAHHIPDVITYSRLRQSVFNCAIQPQWMAATL 300OY 258 AACYNNOQVFKGAVKIRKGQAVTLMDDATNMPPAKAIIYQMEBIYHRIPDSPSSKTR 317
Db 301 AACYNNOQVFKGAVKIRKGQAVTLMDDATNMPPAKAIIYQMEBIYHRIPDSPSSKTR 360OY 318 QIISTIRTQNLPCNQLSRSHYSPYIYSPVFLAALSQWLTLSQTYEDYQOTGEH 374
Db 361 QIISTIRTQNLPCNQLSRSHYSPYIYSPVFLAALSQWLTLSQTYEDYQOTGEH 417

RESULT 6

ID ADB58261 standard; protein; 417 AA.

AC ADB58261;

XX DT 29-JAN-2004 (first entry)

XX DE Human Protein P37268, SEQ ID NO 4132.

XX KW Human; pain; neuronal tissue; gene therapy; spinal segmental nerve injury; chronic constriction injury; CCI; spared nerve injury; SNI; Chung.

XX OS Homo sapiens.

XX PN W02003016475-A2.

XX PD 2-FEB-2003.

XX PR 14-AUG-2002; 2002WO-US025765.

PR 14-AUG-2001; 2001US-0312147P.

PR 01-NOV-2001; 2001US-0316382P.

PR 26-JUL-2001; 2001US-0333347P.

XX PA (GENO) GEN HOSPITAL CORP.

PA (FARB) BAYER AG.

XX PI Wolf C, Durso D, Beaufort K, Costigan M;

XX DR WPI; 2003-125312/25.

XX DR GENBANK; P37268.

XX PT New composition comprising two or more isolated polypeptides, useful for preparing a medicament for treating pain in an animal.

PS XX Claim 1; Page; 1017pp; English.

CC CC The invention discloses a composition comprising two or more isolated rat or human polynucleotides or a polynucleotide which represents a fragment, derivative or allelic variation of the nucleic acid sequence. Also claimed are a vector comprising the novel polynucleotide, a host cell comprising the vector, a method for identifying a nucleotide sequence which is differentially regulated in an animal subjected to pain and a kit to perform the method, an array, a method for identifying an agent that increases or decreases the expression of the polynucleotide sequence that is differentially expressed in neuronal tissue of a first animal subjected to pain, a method for identifying a compound which regulates the expression of a polynucleotide sequence which is differentially expressed in an animal subjected to pain, a method for identifying a compound that regulates the activity of one or more of the polynucleotides, a method for producing a pharmaceutical composition, a method for identifying a compound or small molecule that regulates the activity in an animal of one or more of the polypeptides given in the specification, a method for identifying a compound useful in treating pain and a pharmaceutical composition comprising the one or more polypeptides or their antibodies. The polynucleotide or the compound that modulates its activity is useful for preparing a medicament for treating pain (e.g. spinal segmental nerve injury (SNI) (in an animal (e.g. gene therapy). The sequence presented is a human protein (shown in Table 2 of the specification) which is differentially expressed during pain. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic form directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.

CC CC

XX PT

DE Human Protein P37268, SEQ ID NO 4136.
 XX |||||
 KW Human; pain; neuronal tissue; gene therapy;
 KW spinal segmental nerve injury; chronic constriction injury; CCI;
 KW spared nerve injury; SNI; Chung.
 XX Homo sapiens.
 OS WO2003016475-A2.
 XX PD 27-FEB-2003.
 XX PF 14-AUG-2002; 2002WO-US025765.
 XX PR 14-AUG-2001; 2001US-0312147P.
 PR 01-NOV-2001; 2001US-0346382P.
 PR 26-NOV-2001; 2001US-0333347P.
 XX PA (GEHO) GEN HOSPITAL CORP.
 PA (FARB) BAYER AG.
 XX PI Wolf C, D'urso D, Befort K, Costigan M;
 DR WPI; 2003-268312/26.
 XX DR GENBANK; P37268.
 PT New composition comprising two or more isolated polypeptides, useful for
 preparing a medicament for treating pain in an animal.
 XX PS Claim 1; Page; 1017pp; English.
 XX The invention discloses a composition comprising two or more isolated rat
 CC or human polynucleotides or a polynucleotide which represents a fragment,
 CC derivative or allelic variation of the nucleic acid sequence. Also
 CC claimed are a vector comprising the novel polynucleotide, a host cell
 CC comprising a vector, a method for identifying a nucleotide sequence
 CC which is differentially regulated in an animal subjected to pain and a
 CC kit to perform the method, an array, a method for identifying an agent
 CC that increases or decreases the expression of the polynucleotide sequence
 CC that is differentially expressed in neuronal tissue of a first animal
 CC subjected to pain, a method for identifying a compound which regulates
 CC the expression of a polynucleotide sequence which is differentially
 CC expressed in an animal subjected to pain, a method for identifying a
 CC compound that regulates the activity of one or more of the
 CC polynucleotides, a method for producing a pharmaceutical composition, a
 CC method for identifying a compound or small molecule that regulates the
 CC activity in an animal of one or more of the polypeptides given in the
 CC specification, a method for identifying a compound useful in treating
 CC pain and a pharmaceutical composition comprising the one or more
 CC polypeptides or their antibodies. The polynucleotide or the compound that
 CC modulates its activity is useful for preparing a medicament for treating
 CC pain (e.g. spinal segmental nerve injury (Chung), chronic constriction
 CC injury (CCI) and spared nerve injury (SNI) in an animal (e.g. gene
 CC therapy). The sequence presented is a human protein (shown in Table 2 of
 CC the specification) which is differentially expressed during pain. Note:
 CC the sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic form directly from WIPO at
 CC [fp.wipo.int/pub/published_pct_sequences](http://wipo.int/pub/published_pct_sequences).
 XX SQ Sequence 417 AA;

Query Match 98.4%; Score 1920.5; DB 7; Length 417;
 Best Local Similarity 89.7%; Pred. No. 6.1e-192;
 Matches 374; Conservative 0; Mismatches 0; Indels 43; Gaps 1;

OY 1 MEFYVKCLGHPPEEFVNLVRIGERIGKRVKMPMDMDSLSLSSKTYKLYNQTSSSFAAVIQQA 60
 1 MEFYVKCLGHPPEEFVNLVRIGERIGKRVKMPMDMDSLSLSSKTYKLYNQTSSSFAAVIQQA 60
 Db 61 LPDEMRNNAVCIFVYLRAALDTEDMTSVKEKPLLNPHSTLYQDPWRFMSKEDKDRQ 120
 Oy 61 LPDEMRNNAVCIFVYLRAALDTEDMTSVKEKPLLNPHSTLYQDPWRFMSKEDKDRQ 120
 Db 61 LDGEMRNNAVCIFVYLRAALDTEDMTSVKEKPLLNPHSTLYQDPWRFMSKEDKDRQ 120

QY 121 VLEDFPM-----YCHYAGLVG 137
 Db 121 VLEDFPM-----YCHYAGLVG 180
 QY 138 IGLSRFLPSAASRFEDPLVGEDPTERANSMGMLQKTNITRDYLEDQGGRWPOEWSRY 197
 Db 181 IGLSLRLPSAASRFEDPLVGEDPTERANSMGMLQKTNITRDYLEDQGGRWPOEWSRY 240
 QY 198 KKLGDPAKPNENIDLAQCLMLITNALHHPDVITLSPRNRNQSFVNFCAPQWMAITL 257
 Db 241 KKLGDPAKPNENIDLAQCLMLITNALHHPDVITLSPRNRNQSFVNFCAPQWMAITL 300
 QY 258 AACYNHQOVFKGAVKIRKGQAVTLMDDATNNPAVAKIYQMEYHRIDSDPSSKTR 317
 Db 301 AACYNHQOVFKGAVKIRKGQAVTLMDDATNNPAVAKIYQMEYHRIDSDPSSKTR 360
 QY 318 QIQRSTQNLPNCOLSRSHYSPTIYSPFLNMILALAISWQLTLSQYTEDYVOTGH 374
 Db 361 QIQRSTQNLPNCOLSRSHYSPTIYSPFLNMILALAISWQLTLSQYTEDYVOTGH 417

RESULT 8
 AD658273
 ID ADE58273 Standard; protein; 417 AA.
 XX AC ADE58273;
 XX DT 29-JAN-2004 (first entry)
 XX DE Human Protein P37268, SEQ ID NO 4144.
 XX OS Homo sapiens.
 XX PN WO2003016475-A2.
 XX PR 14-AUG-2002; 2002WO-US025765.
 XX PD 27-FEB-2003.
 XX PA (GEHO) GEN HOSPITAL CORP.
 XX PA (FARB) BAYER AG.
 XX PI Wolf C, D'urso D, Befort K, Costigan M;
 DR WPI; 2003-268312/26.
 XX DR GENBANK; P37268.
 PT New composition comprising two or more isolated polypeptides, useful for
 preparing a medicament for treating pain in an animal.
 XX PS Claim 1; Page; 1017pp; English.
 XX The invention discloses a composition comprising two or more isolated rat
 CC or human polynucleotides or a polynucleotide which represents a fragment,
 CC derivative or allelic variation of the nucleic acid sequence. Also
 CC claimed are a vector comprising the novel polynucleotide, a host cell
 CC comprising a vector, a method for identifying a nucleotide sequence
 CC which is differentially regulated in an animal subjected to pain and a
 CC kit to perform the method, an array, a method for identifying an agent
 CC that increases or decreases the expression of the polynucleotide sequence
 CC that is differentially expressed in neuronal tissue of a first animal
 CC subjected to pain, a method for identifying a compound which regulates
 CC the expression of a polynucleotide sequence which is differentially
 CC expressed in an animal subjected to pain, a method for identifying a
 CC compound that regulates the activity of one or more of the

CC polynucleotides, a method for producing a pharmaceutical composition, a
 CC method for identifying a compound or small molecule that regulates the
 CC activity in an animal or one or more of the polypeptides given in the
 CC specification, a method for identifying a compound useful in treating
 CC pain and a pharmaceutical composition comprising the one or more
 CC polypeptides or their antibodies. The polynucleotide or the compound that
 modulates its activity is useful for preparing a medicament for treating
 pain (e.g. spinal segmental nerve injury (Chung), chronic constriction
 injury (CCI) and spared nerve injury (SNI) in an animal (e.g. gene
 therapy). The sequence presented is a human protein (shown in Table 2 of
 the specification) which is differentially expressed during pain. Note:
 CC The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic form directly from WIPO at
 CC ftp://wipo.int/pub/published_pct_sequences.
 XX SQ Sequence 417 AA:

Query Match	98.4%	Score 1920.5; DB 7; Length 417;
Best Local Similarity	89.7%	Pred. No. 6.1e-192;
Matches	374;	Mismatches 0; Indels 43; Gaps 1;

OY 1 MEFVKCLGHPPEEFLNVLVRIGRKWRKPKMDQDLSLSSIKTYKLYNQTSRFAVTOA 60
 Db 1 MEFVKCLGHPPEEFLNVLVRIGRKWRKPKMDQDLSLSSIKTYKLYNQTSRFAVTOA 60
 OY 61 LDGEMRNACIVFYLVRALDTEDMTSVEKVKPLHNFSFLYQDPWRFMSEKDRQ 120
 Db 61 LDGEMRNACIVFYLVRALDTEDMTSVEKVKPLHNFSFLYQDPWRFMSEKDRQ 120
 OY 121 VDEPPT----- 137
 Db 121 VLDGFPTISLERPLNLAEKKVQTIVADICRRMGICMAEFLDKHVTSEQENDKYCHVAGLV 180
 OY 138 IGSRLFSASEREDPLVGDTERANSWMGLFQLOKNTNIROYLEDOQGGRBFWPQEWSRYV 197
 Db 181 IGSLRFLSASEFEDPLVGDTERANSWMGLFQLOKNTNIROYLEDOQGGRBFWPQEWSRYV 240
 OY 198 KKGDFAKENPDILAVOCINELITNALHHIPDVITYLRLNRNSVENCAIPOMATL 257
 Db 241 KKLGDFAKENPDILAVOCINELITNALHHIPDVITYLRLNRNSVENCAIPOMATL 300
 OY 258 AACCNQQQPKGAKIRKGAVTJMDATMPAKAIQYMERIYHLPDSPSSKTR 317
 Db 301 AACCTNNQQPKGAKIRKGAVTJMDATMPAKAIQYMERIYHLPDSPSSKTR 360
 OY 318 QISTIRTQNLPCOLISRSRHSYSPYLFSWMLALSWOYLTSQVTEDYVOTGEH 374
 Db 361 QISTIRTQNLPCOLISRSRHSYSPYLFSWMLALSWOYLTSQVTEDYVOTGEH 417

RESULT 9
 ADJ94872 ID ADJ94872 standard; protein; 417 AA.
 XX AC ADJ94872;
 XX DT 06-MAY-2004 (first entry)
 XX DE Novel NMR protein sequence #50.

XX antidiabetic; anorectic; cardiotonic; hypotensive; antiarteriosclerotic;
 KW anorectic; viricide; antibacterial; fungicide; protozoacide; nootropic;
 KW neuroprotective; anticarcinonian; anticonvulsant; osteopathic;
 KW antiarthritic; antiinflammatory; dermatological; antiasthmatic;
 KW antilipemic; gene therapy; metabolic disorder; diabetes; obesity;
 KW infections disease; anorexia; cardiovascular disease;
 KW hypertension; atherosclerosis; cancer; neurodegenerative disorder;
 KW Alzheimer's disease; Parkinson's disease; epilepsy; immune disorder;
 KW osteoarthritis; hematopoietic disorder; inflammatory skin disorder;
 KW asthma; dyslipidemia; neurogenesis; cell differentiation;
 KW cell proliferation; hematopoiesis; wound healing; angiogenesis;
 KW chromobone mapping; tissue typing; pharmacogenomic.
 XX

OS Homo sapiens.	
XX	
PN WO2003040325-A2.	
XX	
PD 15-MAY-2003.	
XX	
PR 05-NOV-2002; 2002WO-US035464.	
XX	
PR 05-NOV-2001; 2001US-0338526P.	
PR 06-NOV-2001; 2001US-033972P.	
PR 09-NOV-2001; 2001US-0348583P.	
PR 15-NOV-2001; 2001US-0335610P.	
PR 16-NOV-2001; 2001US-033843P.	
PR 20-NOV-2001; 2001US-0331630P.	
PR 21-NOV-2001; 2001US-0332152P.	
PR 27-NOV-2001; 2001US-0333461P.	
PR 28-NOV-2001; 2001US-0333912P.	
PR 29-NOV-2001; 2001US-033400P.	
PR 30-NOV-2001; 2001US-033421P.	
PR 04-DEC-2001; 2001US-0334526P.	
PR 04-DEC-2001; 2001US-033656P.	
PR 07-DEC-2001; 2001US-0338214P.	
PR 07-DEC-2001; 2001US-0338350P.	
PR 10-DEC-2001; 2001US-0339006P.	
PR 10-DEC-2001; 2001US-0339008P.	
PR 11-DEC-2001; 2001US-0339286P.	
PR 01-FEB-2002; 2002US-035380P.	
PR 01-FEB-2002; 2002US-035388P.	
PR 04-FEB-2002; 2002US-035432P.	
PR 04-FEB-2002; 2002US-0354593P.	
PR 05-MAR-2002; 2002US-0361925P.	
PR 05-MAR-2002; 2002US-036220P.	
PR 05-MAR-2002; 2002US-0360148P.	
PR 07-FEB-2002; 2002US-0361870P.	
PR 05-MAR-2002; 2002US-0361770P.	
PR 05-MAR-2002; 2002US-0361833P.	
PR 13-MAR-2002; 2002US-036197P.	
PR 13-MAR-2002; 2002US-0364227P.	
PR 17-MAY-2002; 2002US-0381621P.	
PR 28-MAY-2002; 2002US-0383675P.	
PR 17-JUL-2002; 2002US-0396703P.	
PR 06-AUG-2002; 2002US-0401522P.	
PR 07-AUG-2002; 2002US-0401787P.	
PR 15-AUG-2002; 2002US-0403619P.	
PR 20-AUG-2002; 2002US-0404821P.	
PR 23-AUG-2002; 2002US-0405368P.	
PR 23-AUG-2002; 2002US-0405402P.	
PR 23-AUG-2002; 2002US-0405496P.	
PR 23-AUG-2002; 2002US-0405631P.	
PR 26-AUG-2002; 2002US-0406125P.	
PR 04-NOV-2002; 2002US-00287226.	
XX (CURA-) CURAGEN CORP.	
PI Agee ML, Alsbrook JP, Berghs C, Boldog FL, Burgess CE, Chant JS;	
PI Chaudhuri A, Dipippo VA, Edinger SR, Eisen A, Elerman K, Ganguli EA, Goracci L, Malayankar UM, Macdougall JR, Mees PS, Miller CE, Millet I;	
PI Ooi CE, Ort T, Padiaru M, Paturrajan M, Rabatelli L, Riger DK;	
PI Rothenberg ME, Shenvoy SG, Spaderna SK, Spytek KA, Taupier RJ;	
PI Vernet GM, Zerhouni BD, Zhong M;	
XX DR WPI; 2003-441551-41.	
DR N-PSDB; ADJ94871.	

XX
PT New isolated NOVX polypeptides and polynucleotides, useful for
PT preventing, diagnosing or treating NOVX-associated disorders, e.g.
PT osteoarthritis, obesity, atherosclerosis, cancer, Parkinson's disease,
PT asthma, or infections.
XX
PS Claim 1; SEQ ID NO 100; 80pp; English.

CC The invention relates to novel isolated polypeptides, mature forms of
CC these, or a sequence that is at least 95 % identical to, or having one or
CC more conservative amino acid substitutions in the polypeptides. The
CC manufacture of a medicament for treating a syndrome associated with a
CC human disease, preferably a NOVX-associated disorder. The nucleic acid
CC molecules, polypeptides and antibodies are useful in the
CC prevention, or diagnosing diseases such metabolic disorders, diabetes,
CC obesity, infectious diseases (viral, bacterial, fungal, helminthic, and
CC protozoal), anorexia, cancer, cardiovascular diseases (hypertension,
CC atherosclerosis), neurodegenerative disorders, Alzheimer's disease,
CC parkinson's disease, epilepsy, immune disorders (osteoarthritis'), and various
CC hematopoietic disorders, inflammatory skin disorders, asthma, and various
CC dyslipidemias. The nucleic acids and polypeptides may also be used as
CC targets for the identification of small molecules that modulate or
CC inhibit e.g. neurogenesis, cell differentiation, cell proliferation,
CC hematopoiesis, wound healing and angiogenesis, in gene therapy, in
CC generation of antibodies that bind immunospecifically to NOVX substances
CC further used as hybridization probes in chromosome mapping, tissue
CC typing, preventive medicine, and pharmacogenomics. This sequence
XX corresponds to one of the NOVX polypeptides of the invention.

SQ Sequence 417 AA;

Query Match 98.4%; Score 1920.5; DB 7; Length 417;
Best Local Similarity 89.7%; Pred. No. 6.1e-192; Mismatches 0; Indels 43; Gaps 1;
Matches 374; Conservative 0; OS Homo sapiens.

1 MEFVKCLGHPEPFYNUVRIGGKRKMPKDQDSSLSKTCYKLNQTSRSFAVIOA 60
1 MEFVKCLGHPEPFYNUVRIGGKRKMPKDQDSSLSKTCYKLNQTSRSFAVIOA 60

QY 61 LDGEMNAVCIFYLVLRALEDDMTISVEKKVPLHNHFSFLYQDPDWRFMESKEKDRQ 120
61 LDGEMNAVCIFYLVLRALEDDMTISVEKKVPLHNHFSFLYQDPDWRFMESKEKDRQ 120

Db 61 LDGEMNAVCIFYLVLRALEDDMTISVEKKVPLHNHFSFLYQDPDWRFMESKEKDRQ 120

QY 121 VLEDFTP-----YCHYVAGLV 137
121 VLEDFTP-----YCHYVAGLV 137

Db 121 VLEDFTP-----YCHYVAGLV 137

QY 121 VLEDFTP-----YCHYVAGLV 137
121 VLEDFTP-----YCHYVAGLV 137

Db 121 VLEDFTP-----YCHYVAGLV 137

XX

CC Sequence 417 AA;

Query Match 98.4%; Score 1920.5; DB 8; Length 417;
Best Local Similarity 89.7%; Pred. No. 6.1e-192; Mismatches 0; Indels 43; Gaps 1;
Matches 374; Conservative 0; OS Homo sapiens.

1 MEFVKCLGHPEPFYNUVRIGGKRKMPKDQDSSLSKTCYKLNQTSRSFAVIOA 60
1 MEFVKCLGHPEPFYNUVRIGGKRKMPKDQDSSLSKTCYKLNQTSRSFAVIOA 60

QY 61 LDGEMNAVCIFYLVLRALEDDMTISVEKKVPLHNHFSFLYQDPDWRFMESKEKDRQ 120
61 LDGEMNAVCIFYLVLRALEDDMTISVEKKVPLHNHFSFLYQDPDWRFMESKEKDRQ 120

Db 61 LDGEMNAVCIFYLVLRALEDDMTISVEKKVPLHNHFSFLYQDPDWRFMESKEKDRQ 120

QY 121 VLEDFTP-----YCHYVAGLV 137
121 VLEDFTP-----YCHYVAGLV 137

Db 121 VLEDFTP-----YCHYVAGLV 137

QY 121 VLEDFTP-----YCHYVAGLV 137
121 VLEDFTP-----YCHYVAGLV 137

Db 121 VLEDFTP-----YCHYVAGLV 137

XX

Oy 138 IGLSRLFSASEFDPFLYGEDTERANSMGLFLOKTNITRDYLEDQDQGREFWQEVNSRYV 197
 CC ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Db 181 IGLSRLFSASEFDPFLYGEDTERANSMGLFLOKTNITRDYLEDQDQGREFWQEVNSRYV 240
 CC or unwanted expression of the protein.
 CC human Farnesyl-diisophsate farnesyltransferase 1 (squalene synthase)
 CC used in a sequence comparison with the novel human enzyme of the
 CC invention.

Oy 198 KKLGDFAKPENDLAVOCNLNEITNALHHIPVITYLSRQLNSVFCAIPOMAIALT 257
 CC ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Db 241 KKLGDFAKPENDLAVOCNLNEITNALHHIPVITYLSRQLNSVFCAIPOMAIALT 300
 CC ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 XX RESULT 11
 ID ADN96864 standard; protein; 417 AA.
 XX AC ADN96864;
 XX DT 26-AUG-2004 (first entry)
 DE Farnesyl-diisophsate farnesyltransferase 1 (squalene synthase).
 KW disease diagnosis; gene expression associated disorder; gene expression;
 KW enzyme peptide; human; enzyme; farnesyl-diisophsate farnesyltransferase 1; squalene synthase.
 KW farnesyl-diisophsate farnesyltransferase 1; squalene synthase.
 OS Homo sapiens.
 PN US2004106179-A1.
 XX PD 03-JUN-2004.
 PP 20-AUG-2003; 2003US-00644021.
 PR 29-MAR-2001; 2001US-00820004.
 XX PA (APPL-) APPLERA CORP.
 XX PI Wei M, Yan C, Di Francesco V, Beasley E;
 DR WPI; 2004-419461/39.
 XX PT New isolated enzyme proteins, useful for diagnosing or treating diseases
 PT characterized by absence, inappropriate, or unwanted expression of the
 PT protein, or as a reagent, in assays for determining levels of protein in
 PT biological sample.

PS Disclosure; SEQ ID NO 4; 76pp; English.

XX The invention describes an isolated enzyme peptide (1) comprising a
 CC defined sequence of 374 amino acids. Also described are: an isolated
 CC antibody that selectively binds to (1); a method for producing the
 CC peptides; a method for detecting the presence of the peptides; a method
 CC for identifying a modulator of the peptide; a method for identifying an
 CC agent that binds to the peptide; a pharmaceutical composition comprising
 CC an agent identified by the method of (5) and a pharmaceutical carrier; a
 CC method for treating a disease or condition mediated by a human enzyme;
 CC a method for identifying a modulator of the expression of the
 CC peptide; and an isolated human enzyme peptide having an amino acid
 CC sequence that shares at least 70-90% homology with SEQ ID NO. 2.
 CC Specifically claimed is an enzyme peptide comprising 374 amino acids (SEQ
 CC ID NO. 2). The peptides are useful for substantial or specific assays,
 CC e.g. biological, or drug screening assays; as a reagent in assays for
 CC determining levels of protein in biological sample; and as markers for
 CC tissues where the corresponding protein is expressed. It can also be used
 CC to screen a compound for the ability to stimulate or inhibit interaction
 CC between the enzyme protein and a molecule that normally interacts with
 CC the enzyme protein. They area also useful as a target for diagnosing a

CC disease or predisposition to disease mediated by the peptide. It can also
 CC be used for treating disorders characterised by absence, inappropriate,
 CC or unwanted expression of the protein. This is the amino acid sequence
 CC human Farnesyl-diisophsate farnesyltransferase 1 (squalene synthase)
 CC used in a sequence comparison with the novel human enzyme of the
 CC invention.

SQ Sequence 417 AA;

Query Match	98.4%	Score	1920.5	DB	8	Length	417
Best Local Similarity	89.7%	Pred.	No.	6.1e-	-192	Mismatches	0
Matches	374	Conservative	0	Indels	43	Gaps	1

Db 318 QIISTIRTONLPCNQLTSRSHSPIVLSFVMILALASWQYLTLSQVTDYQVTGEH 374
 CC ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Db 361 QIISTIRTONLPCNQLTSRSHSPIVLSFVMILALASWQYLTLSQVTDYQVTGEH 417
 CC ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

Oy 61 LGEMRMVACIVTYLVRALDTEDDMTISVERKVLPHNFSFLYOPDWRFMESKEKDQ 120
 CC ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Db 61 LGEMRMVACIVTYLVRALDTEDDMTISVERKVLPHNFSFLYOPDWRFMESKEKDQ 120
 CC ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Oy 121 VIADPFT-----
 CC ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Db 121 VIADPFT-----
 CC ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Oy 138 IGLSRLFSASEFDPFLYGEDTERANSMGLFLOKTNITRDYLEDQDQGREFWQEVNSRYV 197
 CC ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Db 181 IGLSRLFSASEFDPFLYGEDTERANSMGLFLOKTNITRDYLEDQDQGREFWQEVNSRYV 240
 CC ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Oy 198 KKLGDFAKPENDLAVOCNLNEITNALHHIPVITYLSRQLNSVFCAIPOMAIALT 257
 CC ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Db 241 KKLGDFAKPENDLAVOCNLNEITNALHHIPVITYLSRQLNSVFCAIPOMAIALT 300
 CC ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
 XX RESULT 12
 ID ADT79957
 XX AC ADT79957 standard; protein; 417 AA.
 XX DT 16-DEC-2004 (first entry)
 DE Human squalene synthase protein #1.
 XX PN US2004102405-A1.
 XX PD 27-MAY-2004.
 XX PR 23-NOV-2002; 2002US-00304125.
 XX PR 23-NOV-2002; 2002US-00304125.
 XX PR (ISTS-) ISIS PHARM INC.
 XX PI Freier SM, Bennett CF, Dean NM, Dobbie KW;
 XX DR WPI; 2004-399735/37.
 XX DR N-PDBB; ADT79815.

PT New oligonucleotide targeted to a nucleic acid molecule encoding squalene synthase, useful in diagnosing and treating atherosclerosis.
 XX
 PS Disclosure; Page 23-24; 67pp; English.
 XX
 CC The invention relates to a new compound 8-80 nucleobases in length (an antisense oligonucleotide) targeted to a nucleic acid molecule encoding squalene synthase (also known as farnesyl diphosphate farnesyl transferase 1), where the compound specifically hybridises with the nucleic acid molecule encoding human squalene synthase appearing as ADT9815 and inhibits the expression of squalene synthase. Also included are inhibiting the expression of squalene synthase in cells or tissues, screening for a modulator of squalene synthase, a diagnostic method for identifying a disease state, a kit or assay device comprising the compound and treating an animal having a disease or condition associated with squalene synthase. The compound and methods are useful in diagnosing and treating disorders related to cholesterol biosynthesis e.g. atherosclerosis, coronary heart disease and hypercholesterolaemia. The present sequence is a squalene synthase protein sequence.
 XX
 SQ Sequence 417 AA;

Query Match 98.4%; Score 1920.5; DB 9; Length 417;
 Best Local Similarity 89.7%; Pred. No. 6.1e-192;
 Matches 374; Conservative 0; Mismatches 0; Indels 43; Gaps 1;
 Qy 1 MEVVKCLGHPPEEFYNLVRFRIGGKRKMPMDQDSLSSSLKTYKYLQNTSRFAVIOA 60
 Db 1 METVKCLGHPPEEFYNLVRFRIGGKRKMPMDQDSLSSSLKTYKYLQNTSRFAVIOA 60
 Qy 61 LDGEMRNACIVFYLVRLALDTDDMTSVKEKVLPLNPHSFLYQPDWRFMESKEKDRQ 120
 61 LDGEMRNACIVFYLVRLALDTDDMTSVKEKVLPLNPHSFLYQPDWRFMESKEKDRQ 120
 Qy 121 VLEDRAFT-----YCHYVAGLV 137
 121 VLEDRAFT-----YCHYVAGLV 137
 Db 198 KKLGDFAKPENIDLAVOCNLITNALHHPDVTYLSRLRNGSVNFCAPQVMAIATL 257
 241 KKLGDFAKPENIDLAVOCNLITNALHHPDVTYLSRLRNGSVNFCAPQVMAIATL 300
 Qy 258 AACYNQQVFKGAVKIRKGQAVALMDATNPAPKAIIYQMEIYHRIPDSSSKTR 317
 301 AACYNQQVFKGAVKIRKGQAVALMDATNPAPKAIIYQMEIYHRIPDSSSKTR 360
 Db 318 QIISTIRTONLPNCOLISRSRHSYSPIVLSFVMLAALSQWQLTLSQVTDYVQTGEH 374
 361 QIISTIRTONLPNCOLISRSRHSYSPIVLSFVMLAALSQWQLTLSQVTDYVQTGEH 417
 RESULT 13
 ADY16510
 ID ADY16510 standard; protein; 417 AA.
 XX
 AC ADY16510;
 XX
 DT 05-MAY-2005 (first entry)
 XX
 DE PRO polypeptide SEQ ID NO 2316.

Query Match 98.4%; Score 1920.5; DB 9; Length 417;
 Best Local Similarity 89.7%; Pred. No. 6.1e-192;
 Matches 374; Conservative 0; Mismatches 0; Indels 43; Gaps 1;
 Qy 1 MEVVKCLGHPPEEFYNLVRFRIGGKRKMPMDQDSLSSSLKTYKYLQNTSRFAVIOA 60
 Db 1 METVKCLGHPPEEFYNLVRFRIGGKRKMPMDQDSLSSSLKTYKYLQNTSRFAVIOA 60
 Qy 61 LDGEMRNACIVFYLVRLALDTDDMTSVKEKVLPLNPHSFLYQPDWRFMESKEKDRQ 120
 61 LDGEMRNACIVFYLVRLALDTDDMTSVKEKVLPLNPHSFLYQPDWRFMESKEKDRQ 120
 Qy 121 VLEDRAFT-----YCHYVAGLV 137
 121 VLEDRAFT-----YCHYVAGLV 137
 Db 198 KKLGDFAKPENIDLAVOCNLITNALHHPDVTYLSRLRNGSVNFCAPQVMAIATL 257
 241 KKLGDFAKPENIDLAVOCNLITNALHHPDVTYLSRLRNGSVNFCAPQVMAIATL 300
 Qy 258 AACYNQQVFKGAVKIRKGQAVALMDATNPAPKAIIYQMEIYHRIPDSSSKTR 317
 301 AACYNQQVFKGAVKIRKGQAVALMDATNPAPKAIIYQMEIYHRIPDSSSKTR 360
 Db 318 QIISTIRTONLPNCOLISRSRHSYSPIVLSFVMLAALSQWQLTLSQVTDYVQTGEH 374
 361 QIISTIRTONLPNCOLISRSRHSYSPIVLSFVMLAALSQWQLTLSQVTDYVQTGEH 417
 RESULT 14
 ADZ70390
 ID ADZ70390 standard; protein; 417 AA.
 XX
 AC ADZ70390;
 XX
 DT 30-JUN-2005 (first entry)
 XX
 DE Human protein from lung cancer marker gene RDFT1.
 XX
 OS Homo sapiens.

KW	DNA microarray.
XX	
OS	Homo sapiens.
XX	
PN	WO2005032495-A2.
XX	
PD	14-APR-2005.
XX	
PP	01-OCT-2004; 2004WO-US034163.
XX	
PR	03-OCT-2003; 2003US-0508355P.
XX	
PA	(FARB) BAYER PHARM CORP.
XX	
PT	Taylor I, Pauloski NR, Bigwood D;
XX	
DR	DR NPSDB; ADZT0389.
XX	
CC	Providing a patient diagnosis for lung cancer comprises comparing the level of expression of genes or gene products in a biological sample from the patient with that from a normal individual.
XX	
PS	Claim 3; SEQ ID NO 75; 60pp; English.
XX	
CC	The invention relates to providing a patient diagnosis for lung cancer comprising comparing the level of expression of genes or gene products in a biological sample from the patient with the level of expression of genes or gene products in a biological sample from a normal individual.
CC	Also included are distinguishing between normal and disease tissues, monitoring the response of a patient being treated for lung cancer by administering an anti-cancer agent, identifying a compound useful for the treatment of lung cancer and an array for distinguishing between normal and disease tissues (comprising 2 or more probes corresponding to 2 or more genes selected from any of the 200 nucleotide sequences given in the specification, or 2 or more polypeptides comprising any of the 200 amino acid sequences given in the specification). In providing a patient diagnosis for lung cancer, one or more genes are selected from any of the 200 nucleotide sequences as mentioned in the specification, or one or more gene products are polypeptides selected from any of the 20 amino acid sequences mentioned in the specification. The methods are useful for detecting and treating lung cancer. These may also be used for designing, identifying and optimizing therapeutics for cancer. The present sequence represents a protein from one of the 200 lung cancer marker genes. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp://wipo.int/pub/published_pct_sequences .
XX	
SQ	Sequence 417 AA:
Query Match	98.4%; Score 1920.5; DB 9; Length 417;
Best Local Similarity	89.7%; Pred. No. 6.1e-192; Mismatches 0; Indels 43; Gaps 1;
Matches	374; Conservative 0; MisMatches 0; Indels 43; Gaps 1;
QY	1 MEFVKCLGHPEEFVNLVRFRIGKRKVMKPDQSLSSLKTCYKLNQTSRAVIOA 60
Db	1 MEFVKCLGHPEEFVNLVRFRIGKRKVMKPDQSLSSLKTCYKLNQTSRAVIOA 60
QY	61 LDGEMRNAAVICIFYLVRALDTLEDDMTISVEKKVPLHNPHSFYQOPDRFMESKEKDRQ 120
Db	61 LDGEMRNAAVICIFYLVRALDTLEDDMTISVEKKVPLHNPHSFYQOPDRFMESKEKDRQ 120
QY	121 VLERDPT----- YCHVAGLV 137
Db	121 VLERDPT----- YCHVAGLV 137
QY	138 IGLSRLFSASEFEDPLVGDETERANSMGLFLQOKTNIRDYLEDQGGREFWPOEWSRV 197
Db	181 IGLSRLFSASEFEDPLVGDETERANSMGLFLQOKTNIRDYLEDQGGREFWPOEWSRV 197
QY	198 KKLGPFAKPEPDNLAVQCLQELITLALHHPDVITYLSRNRQSVNFCAIPQMAIATL 257
Db	241 KKLGPFAKPEPDNLAVQCLQELITLALHHPDVITYLSRNRQSVNFCAIPQMAIATL 300
QY	258 AACYNNOQVFKGAVKURKGAVTLMDATNNPAKAIYQMEIYHRIPSPSSKTR 317
Db	301 AACYNNOQVFKGAVKURKGAVTLMDATNNPAKAIYQMEIYHRIPSPSSKTR 360
QY	318 QIISTIRIQNLNCOLISRSRHYSPIVISFWMLAISWQVLTLSQVTEDYVQTGEH 374
Db	361 QIISTIRIQNLNCOLISRSRHYSPIVISFWMLAISWQVLTLSQVTEDYVQTGEH 417
	RESULT 15
	AAR52606 standard; protein; 417 AA.
	AAR52606;
	05-DEC-1994 (first entry)
	DE Human squalene synthase.
	XX Squalene synthase; sterol; metabolism; hypercholesterolemia; atherosclerosis; treatment; therapy; prevention.
	XX Homo sapiens.
	XX GB2272442-A.
	XX PD 18-MAY-1994.
	XX PF 09-NOV-1993; 93GB-00023035.
	XX PR 11-NOV-1992; 92GB-00023610.
	XX PA (ZENE) ZENECA LTD.
	XX PT Charles AD;
	XX DR WPI; 1994-146577/1-B.
	XX N-PSDB; AAQ02598.
	PT New recombinant human squalene synthase - used for screening for inhibitors which can be used in the treatment or prevention of high cholesterol levels.
	XX PS Claim 3; Page 36-38; 53pp; English.
	XX CC Recombinantly produced human squalene synthase may be used as a source of enzyme in a non-sterol metabolising host for enzymatic studies or for the screening of compounds to identify inhibitors. Selective inhibition of human squalene synthase can be used to lower intracellular cholesterol levels and provide improved treatment for, and prevent, hypercholesterolemia and atherosclerosis. Human squalene synthase can also be used for the production of antibodies
	XX Sequence 417 AA:
Query Match	98.1%; Score 1915.5; DB 2; Length 417;
Best Local Similarity	89.4%; Pred. No. 2.1e-191; Mismatches 1; Indels 43; Gaps 1;
Matches	373; Conservative 0; MisMatches 1; Indels 43; Gaps 1;
QY	1 MEFVKCLGHPEEFVNLVRFRIGKRKVMKPDQSLSSLKTCYKLNQTSRAVIOA 60
Db	1 MEFVKCLGHPEEFVNLVRFRIGKRKVMKPDQSLSSLKTCYKLNQTSRAVIOA 60
QY	61 LDGEMRNAAVICIFYLVRALDTLEDDMTISVEKKVPLHNPHSFYQOPDRFMESKEKDRQ 120
Db	61 LDGEMRNAAVICIFYLVRALDTLEDDMTISVEKKVPLHNPHSFYQOPDRFMESKEKDRQ 120
QY	121 VLERDPT----- YCHVAGLV 137
Db	121 VLERDPT----- YCHVAGLV 137
QY	61 LDGEMRNAAVICIFYLVRALDTLEDDMTISVEKKVPLHNPHSFYQOPDRFMESKEKDRQ 120
Db	61 LDGEMRNAAVICIFYLVRALDTLEDDMTISVEKKVPLHNPHSFYQOPDRFMESKEKDRQ 120
QY	138 IGLSRLFSASEFEDPLVGDETERANSMGLFLQOKTNIRDYLEDQGGREFWPOEWSRV 197
Db	181 IGLSRLFSASEFEDPLVGDETERANSMGLFLQOKTNIRDYLEDQGGREFWPOEWSRV 197
QY	198 KKLGPFAKPEPDNLAVQCLQELITLALHHPDVITYLSRNRQSVNFCAIPQMAIATL 257
Db	241 KKLGPFAKPEPDNLAVQCLQELITLALHHPDVITYLSRNRQSVNFCAIPQMAIATL 300
QY	138 IGLSRLFSASEFEDPLVGDETERANSMGLFLQOKTNIRDYLEDQGGREFWPOEWSRV 197

Db |||||IGLSRLSASEFEDPLVGEDTERANSGLFLQWTNTIRDYLEDQQGGRFWRPEEVNSRYV 240
181 |||||IGLSRLSASEFEDPLVGEDTERANSGLFLQWTNTIRDYLEDQQGGRFWRPEEVNSRYV 240
Qy |||||KKGDFAKPENIDLAVCLNELTNALJHHPDVTYSLRQLQSVENFCALPOVMATL 257
|||||KKGDFAKPENIDLAVCLNELTNALJHHPDVTYSLRQLQSVENFCALPOVMATL 257
Db |||||KKGDFAKPENIDLAVCLNELTNALJHHPDVTYSLRQLQSVENFCALPOVMATL 300
241 |||||KKGDFAKPENIDLAVCLNELTNALJHHPDVTYSLRQLQSVENFCALPOVMATL 300
Qy |||||AACYYNNQQVFKGAVKIRKGQAVTLMMODATNPAAKALIYQMEIYRIPDSPSSKTR 317
318 |||||AACYYNNQQVFKGAVKIRKGQAVTLMMODATNPAAKALIYQMEIYRIPDSPSSKTR 360
Db |||||AACYYNNQQVFKGAVKIRKGQAVTLMMODATNPAAKALIYQMEIYRIPDSPSSKTR 360
Qy |||||QISTIRTQNLPNCQLISRSHYSPIVYLSFVMLIAALSQWYUTTLSQVTEDYVQTGEH 374
361 |||||QISTIRTQNLPNCQLISRSHYSPIVYLSFVMLIAALSQWYUTTLSQVTEDYVQTGEH 417
Db |||||QISTIRTQNLPNCQLISRSHYSPIVYLSFVMLIAALSQWYUTTLSQVTEDYVQTGEH 417

Search completed: March 24, 2006, 16:45:49
Job time : 190 secs

RESULT 2
US-08-351-981-6 Application US/08351981
; Sequence 6, Application US/08351981
; Patent No. 589372
GENERAL INFORMATION:
APPLICANT: Robinson, Gordon W
TITLE OF INVENTION: Squalene Synthetase
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: Burton Rodney
STREET: P.O. Box 4000
CITY: Princeton
STATE: New Jersey
COUNTRY: U.S.A.
ZIP: 08543-4000
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/351,981
FILING DATE:
CLASSIFICATION: 435
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: US/07/911,835
ATTORNEY/AGENT INFORMATION:
NAME: Gaul, Timothy J.
REGISTRATION NUMBER: 33,111
REFERENCE/DOCKET NUMBER: DCTa
TELECOMMUNICATION INFORMATION:
TELEPHONE: (609) 252-5901
TELEFAX: (609) 252-4526
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 417 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
; US-08-351-981-6

Query Match 98.4%; Score 1920.5; DB 1; Length 417;
Best Local Similarity 89.7%; Pred. No. 2.7e-197; Indels 0; Gaps 1;
Matches 374; Conservative 0; Mismatches 0; Indels 43; Gaps 1;

QY 1 MEVKCLGHPEEFYLVLRIGGKRKMPKMDQDSLSSLKTYKYLQNLQNSRSFAVIA 60
Db 1 METVKCLGHPEEFYLVLRIGGKRKMPKMDQDSLSSLKTYKYLQNLQNSRSFAVIA 60
QY 61 LDGEMRNAAVICIVLYVLRAALDTLEDDMTISVEKVKPLHNHFSFLYQDPDWRTMESKEKDRQ 120
Db 61 LDGEMRNAAVICIVLYVLRAALDTLEDDMTISVEKVKPLHNHFSFLYQDPDWRTMESKEKDRQ 120
QY 121 VLEDPT----- YCHYVAGLV 137
Db 121 VLEDPTISLEFRNLAEKYQTVIDICRRNGIGMAEFLDKAVTSEQEWDKYCHYVAGLV 180
QY 138 IGSRLFFSAESEPDPLYGEDTERANSNGLFLQKNTIRDYLEDQDGREFPQEWRSVY 197
Db 181 IGLSRLLSAESEPDPLYGEDTERANSNGLFLQKNTIRDYLEDQDGREFPQEWRSVY 240
QY 198 KKLGDFAKENIDLAVOCINELITNALAHIPDVITMSRLRNOVSVNFCATPQWMAIAL 257
Db 241 KKLGDFAKENIDLAVOCINELITNALAHIPDVITMSRLRNOVSVNFCATPQWMAIAL 300
QY 258 AACYNNOQVKGAVKIRKGQAVTLMDATNPMAVKAIYQMEIYHRIPSDPSSKTR 317
Db 301 AACYNNOQVKGAVKIRKGQAVTLMDATNPMAVKAIYQMEIYHRIPSDPSSKTR 360
QY 318 QISTIRTQNLPCQLISRSHSPITVSPVMALLAISWQYLTLSQTYDVTGEH 374
Db 361 QISTIRTQNLPCQLISRSHSPIVLSFVMLAALSQWYLTLSQTYDVTGEH 417

QY 121 VLEDPT----- YCHYVAGLV 137
Db 121 VLEDPTISLEFRNLAEKYQTVIDICRRNGIGMAEFLDKAVTSEQEWDKYCHYVAGLV 180
QY 138 IGSRLFFSAESEPDPLYGEDTERANSNGLFLQKNTIRDYLEDQDGREFPQEWRSVY 197
Db 181 IGLSRLLSAESEPDPLYGEDTERANSNGLFLQKNTIRDYLEDQDGREFPQEWRSVY 240
QY 198 KKLGDFAKENIDLAVOCINELITNALAHIPDVITMSRLRNOVSVNFCATPQWMAIAL 257
Db 241 KKLGDFAKENIDLAVOCINELITNALAHIPDVITMSRLRNOVSVNFCATPQWMAIAL 300
QY 258 AACYNNOQVKGAVKIRKGQAVTLMDATNPMAVKAIYQMEIYHRIPSDPSSKTR 317
Db 301 AACYNNOQVKGAVKIRKGQAVTLMDATNPMAVKAIYQMEIYHRIPSDPSSKTR 360
QY 318 QISTIRTQNLPCQLISRSHSPITVSPVMALLAISWQYLTLSQTYDVTGEH 374
Db 361 QISTIRTQNLPCQLISRSHSPIVLSFVMLAALSQWYLTLSQTYDVTGEH 417

RESULT 3
US-09-820-004-4 Application US/09820004
; Sequence 4, Application US/09820004
; Patent No. 664385
GENERAL INFORMATION:
APPLICANT: WEI, Ming-Hui et al.
TITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEARIC ACID MOLECULES ENCODING HUMAN ENZYME PROTEINS, AND USES
TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN ENZYME PROTEINS, AND USES
TITLE OF INVENTION: THEREOF
FILE REFERENCE: C1001201
CURRENT APPLICATION NUMBER: US/09/820, 004
CURRENT FILING DATE: 2001-03-29
NUMBER OF SEQ ID NOS: 6
SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 5
; SEQ ID NO 4
; LENGTH: 417
; TYPE: PRX
; ORGANISM: Human
; US-09-820-004-4

Query Match 98.4%; Score 1920.5; DB 2; Length 417;
Best Local Similarity 89.7%; Pred. No. 2.7e-197; Indels 0; Gaps 1;
Matches 374; Conservative 0; Mismatches 0; Indels 43; Gaps 1;

QY 1 MEVKCLGHPEEFYLVLRIGGKRKMPKMDQDSLSSLKTYKYLQNLQNSRSFAVIA 60
Db 1 METVKCLGHPEEFYLVLRIGGKRKMPKMDQDSLSSLKTYKYLQNLQNSRSFAVIA 60
QY 61 LDGEMRNAAVICIVLYVLRAALDTLEDDMTISVEKVKPLHNHFSFLYQDPDWRTMESKEKDRQ 120
Db 61 LDGEMRNAAVICIVLYVLRAALDTLEDDMTISVEKVKPLHNHFSFLYQDPDWRTMESKEKDRQ 120
QY 121 VLEDPT----- YCHYVAGLV 137
Db 121 VLEDPTISLEFRNLAEKYQTVIDICRRNGIGMAEFLDKAVTSEQEWDKYCHYVAGLV 180
QY 138 IGSRLFFSAESEPDPLYGEDTERANSNGLFLQKNTIRDYLEDQDGREFPQEWRSVY 197
Db 181 IGLSRLLSAESEPDPLYGEDTERANSNGLFLQKNTIRDYLEDQDGREFPQEWRSVY 240
QY 198 KKLGDFAKENIDLAVOCINELITNALAHIPDVITMSRLRNOVSVNFCATPQWMAIAL 257
Db 241 KKLGDFAKENIDLAVOCINELITNALAHIPDVITMSRLRNOVSVNFCATPQWMAIAL 300
QY 258 AACYNNOQVKGAVKIRKGQAVTLMDATNPMAVKAIYQMEIYHRIPSDPSSKTR 317
Db 301 AACYNNOQVKGAVKIRKGQAVTLMDATNPMAVKAIYQMEIYHRIPSDPSSKTR 360
QY 318 QISTIRTQNLPCQLISRSHSPITVSPVMALLAISWQYLTLSQTYDVTGEH 374
Db 361 QISTIRTQNLPCQLISRSHSPIVLSFVMLAALSQWYLTLSQTYDVTGEH 417

RESULT 4
US-09-820-004-5 Application US/09820004
; Sequence 5, Application US/09820004
; Patent No. 664385
GENERAL INFORMATION:
APPLICANT: WEI, Ming-Hui et al.
TITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEARIC ACID MOLECULES ENCODING HUMAN ENZYME PROTEINS, AND USES
TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN ENZYME PROTEINS, AND USES
TITLE OF INVENTION: THEREOF
FILE REFERENCE: C1001201
CURRENT APPLICATION NUMBER: US/09/820, 004
CURRENT FILING DATE: 2001-03-29
NUMBER OF SEQ ID NOS: 6
SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 5

Copyright (c) 1993 - 2006 Biocceleration Ltd.

Om protein - protein search, using sw model

Run on: March 24, 2006, 16:42:50 ; Search time 233 seconds

Maximum DB seq length: 0
(without alignments)
1112.479 Million cell updates/sec

Title: US-10-644-021A-2

Perfect score: 1952

Sequence: 1 MEFVKCLGHPPEFFNLVRER.....WQVLTILSQVTEDYVQNGH 374

Scoring table: BL05UN62

Gapop 10.0 , Gapext 0.5

Searched: 2166443 seqs, 705528306 residues

Total number of hits satisfying chosen parameter: 2166443

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : UniProt_05.80;*

1: uniprot_sprot;*

2: uniprot_trembl;*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match Length	DB ID	Description
1	1920.5	98.4	417	1 FDFT_HUMAN
2	1920.5	98.4	417	2 061AA1_HUMAN
3	1920.5	98.4	417	2 Q5R6U3_PONY
4	1820.5	93.3	416	1 FDFT_MOUSE
5	1737	89.0	416	1 Q8BYF5_MOUSE
6	1737	89.0	416	1 FDFT_RAT
7	86.7	86.7	416	1 Q5ZKWL_CHICK
8	1527	78.2	418	2 Q9QYTO_MOUSE
9	1507.5	77.2	368	2 Q9QYD0_MOUSE
10	1208.5	61.9	448	2 Q4RZ3_TEETING
11	755	38.7	415	2 Q5Z666_PAR1A
12	754	38.6	416	2 Q5ADRL_DICOT
13	753.5	38.6	403	2 Q22105_ORYZA
14	749.5	38.4	401	2 Q22106_MAIZE
15	746.5	38.2	411	1 FDFT_NICBEE
16	744	38.1	460	1 FDFT_SCHPO
17	737.5	37.8	411	2 Q24149_TOBAC
18	737.5	37.8	413	2 Q6SYC9_ARTAN
19	737.5	37.8	418	2 Q6SYCB_ARTAN
20	736	37.7	413	2 Q6568B_ARATH
21	735.5	37.7	411	2 Q9XJ31_SOLTJA
22	735.5	37.7	413	2 Q84LE3_LOTUS_Japon
23	732	37.5	413	2 Q23110_ARATH
24	731.5	37.5	413	2 Q22107_SOYBEAN
25	731	37.4	410	1 FDFT_ARATH
26	729.5	37.4	415	2 Q4B6E6_PANGI
27	729	37.3	412	2 Q42761_GLYCRRHIZA
28	725.5	37.2	447	2 Q6BT91_DEBHLA
29	723.5	37.1	448	1 FDFT_CINNAM
30	723.5	37.1	448	2 Q59Y35_CANAL
31	721.5	37.0	411	2 Q9XF02_CAPAN

RESULT 1

FDFT_HUMAN STANDARD; PRT: 417 AA.

ID FDFT_HUMAN

AC P37268; Q9GCT0;

DT 01-OCT-1994 (Rel. 30, Created)

DT 01-OCT-1994 (Rel. 30, Last sequence update)

DT 10-MAY-2005 (Rel. 47, Last annotation update)

DE Farnesyldiphosphate farnesyltransferase (EC 2.5.1.21) (Squalene synthetase) (SQS) (SS) (FPP:FPP farnesylytransferase).

DE Name=FDFT1;

GN Homo sapiens (Human)

OS Homo sapiens (Human)

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominidae; Homo.

OX NEBI_TAXID=9606;

RN [1]

RP NUCLEOTIDE SEQUENCE.

RX MEDLINE=93233634; PubMed=8474436;

RA Robinson G.W., Tsay Y.H., Klenzle B.K., Smith-Monroy C.A., Bishop R.W.,

RA "Conservation between human and fungal squalene synthetases: similarities in structure, function, and regulation.";

RT Mol. Cell. Biol. 13:2706-2727(1993).

RN [2]

RP NUCLEOTIDE SEQUENCE.

RC TISSUE-LIVER;

RX MEDLINE=93286128; PubMed=7685352;

RA Jiang G., McKenzie T.L., Conrad D.G., Shechter I.; "Transcriptional regulation by lovastatin and 25-hydroxycholesterol in HepG2 cells and molecular cloning and expression of the cDNA for the human hepatic squalene synthase.," J. Biol. Chem. 268:12818-12824(1993).

RN [3]

RP NUCLEOTIDE SEQUENCE.

RC TISSUE-LIVER;

RX MEDLINE=94122996; PubMed=8294001; DOI=10.1016/0378-1119(93)90462-C;

RA Summers C., Karst F., Charles A.D.; "Cloning, expression and characterisation of the cDNA encoding human hepatic squalene synthase, and its relationship to phytene synthase.," Gene 135:185-192(1993).

RN [4]

RP NUCLEOTIDE SEQUENCE.

RC TISSUE-LIVER;

RX MEDLINE=95168856; PubMed=7864626; DOI=10.1006/abb1.1995.1095;

RA Soitis D.A., McMahon G., Caplan S.L., Dudas D.A., Chamberlin H.A., Vattay A., Dotavio D., Rucher M.L., Engstrom R.G., Cornell-Kennon S.M.; "Expression, purification, and characterization of the human squalene synthase: use of yeast and baculoviral systems.," Arch. Biochem. Biophys. 316:713-723(1995).

RN [5]

RP NUCLEOTIDE SEQUENCE [LARGE SCALE RNA], AND VARIANT ARG-45. TISSUE=Lung, Muscle, and Urinary bladder;

RX MEDLINE=22388257; PubMed=1247932; DOI=10.1073/pnas.242603899;
 RA Straubberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klaunser R.D., Collins F.S., Wagner L., Shrem C.M., Schuler G.D.,
 RA Altshull S.P., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Ditschenko L., Maruska K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Ronald M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
 RA Raba S.S., Loquellano N.A., Peters G.J.J., Abramson R.D., Mullahy S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smialus D.E.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Villalon D., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahey J., Helton E., Ketteman M., Madan A., Rodrigues S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Jackson R.W., Touché J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Schneeck A., Schein J.E.J.M., Marra M.A.,
 RT "Generation and initial analysis of more than 15,000 full-length human
 RT and mouse cDNA sequences.",
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [6]
 RP X-RAY CRYSTALLOGRAPHY (2.15 ANGSTROMS) OF 39-370.
 PubMed=10896663; DOI=10.1074/jbc.M004132200;
 RA Pandit J., Danley D.E., Schulte G.K., Magazzano S., Pauly T.A.,
 RA Hayward C.M., Hananaka E.S., Thompson J.F., Harwood H.J. Jr.,
 RT "Crystal structure of human squalene synthase. A key enzyme in
 RT cholesterol biosynthesis.",
 RL J. Biol. Chem. 275:30610-30617(2000).
 CC CCA -I CATALYTIC ACTIVITY: Presqualene diphosphate + NADPH = diphosphate
 CC CCA + squalene + NADP(+).
 CC CCA -I COFACTOR: Magnesium.
 CC CCA -I PATHWAY: Critical branch point enzyme of isoprenoid and
 CC CCA cholesterol biosynthesis.
 CC CCA -I SUBUNIT: Monomer (By similarity).
 CC CCA -I SUBCELLULAR LOCATION: Integral membrane protein. Endoplasmic
 CC CCA reticulum.
 CC CCA -- SIMILARITY: Belongs to the phytene/squalene synthetase family.
 CC CCA This Swiss-Prot entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use as long as its content is in no way modified and this statement is not
 CC removed.
 CC CCA
 DR EMBL: L06070; AA060582.1; -; mRNA.
 DR EMBL: L06105; AA036645.1; -; mRNA.
 DR EMBL: X69141; CA04886.1; -; mRNA.
 DR EMBL: S76822; AA033404.1; -; mRNA.
 DR EMBL: BC001353; AAH03573.1; -; mRNA.
 DR EMBL: BC009541; AHH09551.1; -; mRNA.
 DR EMBL: BC029491; AAH29491.1; -; mRNA.
 DR EMBL: A45998; A45998.
 DR PIR; 138245; 138245.
 DR PROTEIN; X-ray; A/B/C=35-370.
 DR Ensembl; ENSG00000079559; Homo sapiens.
 DR HGNC; HGNC:3629; FDF1.
 DR H-InvDB; HIX007319; -.
 DR MTM; 184420; -.
 DR GO; GO:0011621; C:integral to membrane; TAS.
 DR GO; GO:0006694; P:sterol biosynthesis; TAS.
 DR InterPro; IPR020260; Sgu/phyt_synthse.
 DR InterPro; IPR006449; Squal_synth.
 DR Pfam; PF00424; SOS_PSY_1.
 DR TIGRFAMS; TIGR01559; squal synth; 1.
 DR PROSITE; PS01044; SQUALEN_PHTOEN_SYN_1; 1.
 DR PROSITE; PS01045; SQUALEN_PHTOEN_SYN_2; 1.
 KW 3D-structure; Cholesterol biosynthesis; Endoplasmic reticulum;
 KW Isoprene biosynthesis; Lipid synthesis; Magnesium; Polymorphism;
 KW Multifunctional enzyme; NADP; Oxidoreductase; Transf erase; Transmembrane.
 KW Steroid biosynthesis; Sterol biosynthesis; Transferase; Transmembrane.

FT	TRANSMEM	284	304	Potential.
FT	TRANSMEM	384	404	Potential.
FT	VARIANT	45	45	K -> R (in dbSNP:1047695).
FT	VARIANT	392	392	/FTId=VAR_011786. L -> P (in dbSNP:1804473). D -> N (in Ref. 4). T -> A (in Ref. 3).
FT	CONFLICT	353	353	/FTId=VAR_011787.
FT	CONFLICT	402	402	
FT	HELIX	39	50	
FT	HELIX	54	59	
FT	TURN	60	60	
FT	TURN	63	63	
FT	HELIX	64	84	
FT	TURN	86	87	
FT	HELIX	90	103	
FT	TURN	104	105	
FT	TURN	107	108	
FT	TURN	117	118	
FT	HELIX	119	123	
FT	TURN	124	124	
FT	HELIX	125	133	
FT	TURN	134	134	
FT	HELIX	137	158	
FT	HELIX	159	159	
FT	TURN	165	175	
FT	HELIX	176	176	
FT	TURN	177	190	
FT	HELIX	191	191	
FT	TURN	195	199	
FT	HELIX	201	218	
FT	TURN	219	219	
FT	HELIX	220	226	
FT	TURN	227	227	
FT	HELIX	233	236	
FT	TURN	237	239	
FT	HELIX	243	247	
FT	HELIX	249	251	
FT	HELIX	252	267	
FT	TURN	268	269	
FT	HELIX	270	278	
FT	TURN	279	279	
FT	HELIX	283	303	
FT	TURN	304	305	
FT	HELIX	307	310	
FT	TURN	311	311	
FT	TURN	328	329	
FT	HELIX	331	348	
FT	TURN	351	352	
FT	TURN	354	355	
FT	HELIX	355	367	
FT	TURN	368	368	
SO	SEQUENCE	417 AA;	48115 MW;	D36C8C8382F827EC CRC64;
Ov	Query Match	98.4%; Score 1920.5; DB 1; Length 417;		
Ov	Best Local Similarity	89.7%; Pred. No. 8.2e-153; Mis matches 0; Indels 43; Gaps 1;		
Ov	Matches	374;		
Ov	1	MEFVKCLGHPSPEFYNLVRFRIGKRWKMPKNDQDSLSLSSIKTCYKLNQTSRSFAVIAQ	60	
Db	1	MEFVKCLGHPSPEFYNLVRFRIGKRWKMPKNDQDSLSLSSIKTCYKLNQTSRSFAVIAQ	60	
Ov	121	VIEDFPF-----YCHYAGLQ	137	
Db	121	VIEDFPFISLFRNLAKYQWIDICRRNGIGMAEFLDKHVTSDEDWDRKYAGLV	180	
Ov	61	LGDEMNAVCFTYVRLAQLDEDMTISYEKKVPLHNHFSFLYQDPWTFMESKEKDQ	120	
Db	61	LGDEMNAVCFTYVRLAQLDEDMTISYEKKVPLHNHFSFLYQDPWTFMESKEKDQ	120	
Ov	138	IGLSRFLSASEREDPLVGEDETERANSGLFLQTKNTIRDYLEDQQGREFWPOEWRSY	197	
Db	181	IGLSRFLSASEREDPLVGEDETERANSGLFLQTKNTIRDYLEDQQGREFWPOEWRSY	240	

RESULT 3
 US-10-524-972-115
 ; Sequence 116, Application US/10524972
 ; Publication No. US20060031963A1
 ; GENERAL INFORMATION:
 ; APPLICANT: CHIRON SPA
 ; APPLICANT: FONTANA Maria Rita
 ; APPLICANT: PIZZA Margherita
 ; APPLICANT: MASIGNANI Vega
 ; APPLICANT: MONACI Elisabetta
 ; APPLICANT: GONOCOCAL PROTEINS AND NUCLEIC ACIDS
 ; TITLE OF INVENTION:
 ; FILE REFERENCE:
 ; CURRENT APPLICATION NUMBER: US/10/467,657
 ; CURRENT FILING DATE: 2003-08-11
 ; PRIORITY FILING DATE: 2001-02-12
 ; NUMBER OF SEQ ID NOS: 9218
 ; SEQ ID NO 5128
 ; LENGTH: 290
 ; TYPE: PRT
 ; ORGANISM: Neisseria gonorrhoeae
 ; US-10-467-657-5128
 ; Query Match 8.2%; Score 159.5; DB 6; Length 290;
 ; Best Local Similarity 23.6%; Pred. No. 4.2e-07;
 ; Matches 78; Conservative 23.6%; Mismatches 106; Indels 107; Gaps 12;
 ; Prior Application Number: DE 102 38 980.2
 ; Prior Filing Date: 2003-08-18
 ; Prior Application Number: DE 102 38 980.2
 ; Prior Filing Date: 2003-08-18
 ; Prior Application Number: DE 102 38 978.0
 ; Prior Filing Date: 2002-08-20
 ; Prior Application Number: DE 102 38 979.9
 ; Prior Filing Date: 2002-08-20
 ; Prior Application Number: DE 102 53 112.9
 ; Prior Filing Date: 2002-11-13
 ; Prior Application Number: DE 102 58 971.2
 ; Prior Filing Date: 2002-12-16
 ; Prior Application Number: DE 102 58 971.2
 ; Prior Filing Date: 2002-12-16
 ; Software: SeqWin9, version 1.04
 ; SEQ ID NO 116
 ; LENGTH: 309
 ; TYPE: PRT
 ; ORGANISM: Erwinia uredovora
 ; US-10-524-972-116
 Query Match 9.3%; Score 181; DB 6; Length 309;
 Best Local Similarity 22.9%; Pred. No. 4.7e-09;
 Matches 66; Conservative 37; Mismatches 107; Indels 78; Gaps 8;
 Qy 51 SRFRAVTOALDGERMNRNAVCFIVLVRALDTIDDMTISVERKVKPLLNFSRFLYQDWR 110
 Db 18 SKSFRATASKLDFAKTRRSVLMILAWCRHCDVDQTLGFQARQPALOT-----PEQR 70
 Qy 111 FMSSEKEKORQV-----LED 124
 Db 71 LMDLEMKTROQAVGQSOMHQAFOEYAWAHDTAPAYAFDHLEFGFANDVREAOYSQDD 130
 RESULT 4
 US-11-096-568A-6786
 ; Sequence 6786, Application US/11096568A
 ; Publication No. US20060048240A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Alexandrov, Nikolai et al.
 ; TITLE OF INVENTION: Sequence-Determined DNA Fragments and Corresponding Polypeptides I
 ; FILE REFERENCE: 2750-1592P052
 ; CURRENT APPLICATION NUMBER: US/11/096,568A
 ; CURRENT FILING DATE: 2005-04-01
 ; NUMBER OF SEQ ID NOS: 34471
 ; SEQ ID NO 6786
 ; LENGTH: 399
 ; TYPE: PRT
 ; ORGANISM: Glycine max
 ; FEATURE: misc feature
 ; LOCATION: (1)..(399)
 Qy 125 FPTYCHYVAGLVIGLSRLFSASRFEDPLVGEDTERANSMGLFLQKNTIRDYLEDQOGG 184
 Db 131 TLYCYHYVAGVWGMQIMGVHD-----NATLDRACDGLAGFLQTMNIDPVIDDAHAG 184
 Qy 185 RFTWQEWWSRYVKLGDFAKPENIDLAVOCNLNLTNALHHIPDVITYLTSRLRNQSVFN 244
 Db 185 RCVLPASMLEHEGLNKENYAAPENRQLSRIARRVQAEPY-----YLS-----ATAG 233
 Qy 245 FCALP--QWMAITLAACYNNOQVFKGATKIRK-GQAVTLMMDATNP 289
 Db 234 LAGULPLRSWAIAVAKQYR----KIGVKEQMGQQWDQROSSTTP 276

GenCore version 5.1.7
Copyright (c) 1993 - 2006 Biocceleration Ltd.

OM protein - protein search, using sw model

Run on:

March 24, 2006, 16:46:05 ; Search time 40 Seconds
(without alignments)

Sequence: 899 627 Million cell updates/sec

Title: US-10-644-021a-2
Perfect score: 1952

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

PIR_801,*
1: pix1,*
2: pix2,*
3: pix3,*
4: pix4,*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No. Score Query Match Length DB ID Description

ALIGNMENTS

RESULT 1

A45998 A45998 farnezyldiphosphate farnezylytransferase (EC 2.5.1.21) - human

N:Alternate name: squalene synthase

C:Species: Homo sapiens (man)

C:Date: 03-Mar-1994 #sequence_revision 18-Nov-1994 #text_change 09-Jul-2004

C:Accession: A45998; K48057

R:Jiang, G.; McKenzie, T.L.; Conrad, D.G.; Shechter, I.

J: Biol. Chem. 268, 12818-12824, 1993

A:Title: Transcriptional regulation by lovastatin and 25-hydroxycholesterol in HepG2 cell

A:Reference: A45998; MUID:93286128; PMID:7685352

A:Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-417 <JIN>

A:Cross-references: UNIPROT:P37268; UNIPARC:UPI000012A505; GB:L06105; NID:9307431; PIDN:1

A:Experimental source: hepatoma cell line HepG2

A:Note: Sequence extracted from NCBI backbone (NCBIN:133625, NCBI:133626)

R:Robinson, G.W.; Tsay, I.H.; Kleinzle, B.K.; Smith-Monroy, C.A.; Bishop, R.W.

Mol. Cell. Biol. 13, 2706-2717, 1993

A:Title: Conservation between human and fungal squalene synthetases: similarities in str

A:Residues: 1-417 <JIN>

A:Reference number: A48057; MUID:93233634; PMID:8474436

A:Status: preliminary

A:Molecule type: nucleic acid

A:Residues: 1-417 <ROB>

A:Cross-references: UNIPARC:UPI000012A505; GB:L06070; NID:9292509; PIDN:AAA60582_1; PIDC:6

A:Note: Sequence extracted from NCBI backbone (NCBIN:129790, NCBI:129791)

C:Superfamily: farnezyldiphosphate farnezylytransferase

C:Keywords: transferase; transmembrane protein

Query Match 98.4%; Score 1920.5; DB 2; Length 417; Best Local Similarity 89.7%; Pred. No. 7.4e-18; Batches 374; Conservative 0; Mismatches 0; Indels 43; Gaps 1;

Matches 1 1 MEFVKCLGPEEFVNVLVRIGRKWKPKMDPSLSSLLKTCYKLNQTSRFAAVICQY

QY 1 MEFVKCLGPEEFVNVLVRIGRKWKPKMDPSLSSLLKTCYKLNQTSRFAAVICQY

Db 1 MEFVKCLGPEEFVNVLVRIGRKWKPKMDPSLSSLLKTCYKLNQTSRFAAVICQY

Db 1 LDGEMRNACVIFMVLRAIDTUEDMTSVEKKVPLHNFHSLYQPDRFMESEKDRQ

QY 61 LDGEMRNACVIFMVLRAIDTUEDMTSVEKKVPLHNFHSLYQPDRFMESEKDRQ

Db 61 LDGEMRNACVIFMVLRAIDTUEDMTSVEKKVPLHNFHSLYQPDRFMESEKDRQ

Db 121 VLEDPPT-----YCHIVAGWG

QY 121 VLEDPPT-----YCHIVAGWG

QY 138 IGSLRLFSSEFPDPLVGDTERANSWMLFLQKNTIRDYLEDQOGGREFWPOEWWSYV

QY 139 IGSLRLFSSEFPDPLVGDTERANSWMLFLQKNTIRDYLEDQOGGREFWPOEWWSYV

Db 181 IGLSLRLFSSEFPDPLVGDTERANSWMLFLQKNTIRDYLEDQOGGREFWPOEWWSYV

QY 198 KKLGDFAKPENIDLVQCLNELTNALHHPDVITIUSRLRNGSVFNFCFAIPQVMAIAI

QY 199 KKLGDFAKPENIDLVQCLNELTNALHHPDVITIUSRLRNGSVFNFCFAIPQVMAIAI

GenCore version 5.1.7
Copyright (c) 1993 - 2006 Biocceleration Ltd.

Om protein - protein search, using Bw model

Run on:

March 24, 2006, 17:01:21 ; Search time 166 Seconds

(without alignments)
941.375 Million cell updates/sec

Title: US-10-644-021a-2

Perfect score: 1952

Sequence: 1 MEFVKCLIGHPEEEFVNLYRFRIGSKRKWPKMMDPSLSSSLKTCYKLNQTSRSFAAVQIA 60

Scoring table: BL051M62 Gapop 10.0 , Gapext 0.5

Searched: 1867569 seqs, 417829326 residues

Total number of hits satisfying chosen parameters: 1867569

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database : Published Applications AA_Main:**

1: /cgn2_6_ptodata/1/pubaa/US07_PUBCOMB.pep:**

2: /cgn2_5_ptodata/1/pubaa/US08_PUBCOMB.pep:**

3: /cgn2_6_ptodata/1/pubaa/US09_PUBCOMB.pep:**

4: /cgn2_5_ptodata/1/pubaa/US10_PUBCOMB.pep:**

5: /cgn2_5_ptodata/1/pubaa/US11_PUBCOMB.pep:**

6: No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match Length	DB ID	Description
1	1952	100.0	374	3 US-09-820-004-2
2	1952	100.0	374	4 US-10-644-021a-2
3	1952	98.4	417	3 US-09-820-004-4
4	1920.5	98.4	417	3 US-09-820-004-5
5	1920.5	98.4	417	4 US-10-287-100
6	1920.5	98.4	417	4 US-10-644-021a-4
7	1920.5	98.4	417	4 US-10-644-021a-5
8	1915.5	98.1	417	3 US-09-820-004-6
9	1915.5	98.1	417	4 US-10-644-021a-6
10	1692	86.7	416	4 US-10-205-194-67
11	770.5	39.5	455	4 US-10-425-114-66845
12	759.5	38.9	403	4 US-10-425-115-221234
13	753.5	38.6	403	4 US-10-437-963-187558
14	747.5	38.3	401	4 US-10-425-115-222973
15	744	38.1	401	4 US-10-369-493-2381
16	735.5	37.7	428	4 US-10-425-114-66845
17	732	37.5	404	5 US-10-739-930-10172
18	731	37.4	410	4 US-10-024-130A-2
19	723.5	37.1	448	4 US-10-032-585-7355
20	717.5	36.8	413	4 US-10-429-949-5
21	716.5	36.7	408	4 US-10-024-130A-6
22	705.5	35.2	328	4 US-10-424-599-255117
23	695.5	35.6	328	4 US-10-128-714-8505
24	688.5	35.3	444	4 US-10-369-493-22063
25	685.5	35.1	520	4 US-10-128-714-3505
26	676	34.6	354	4 US-10-369-493-13039
27	660.5	33.8	388	4 US-10-425-114-68093

RESULT 1
US-09-820-004-2
; Sequence 2, Application US/09820004
; Patent No. US20020142418A1
; GENERAL INFORMATION:
; APPLICANT: WEI, Ming-Hui et al.
; TITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEIC ACID MOLECULES ENCODING HUMAN ENZYME PROTEINS, AND USES
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: CI001201
; CURRENT APPLICATION NUMBER: US/09/820, 004
; CURRENT FILING DATE: 2001-03-29
; NUMBER OF SEQ ID NOS: 6
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 374
; TYPE: PRT
; ORGANISM: Human
US-09-820-004-2

ALIGNMENTS

Query	Match	Length	DB	ID	Description	Score	DB Score	Length	DB Length	Start	End	Sequence
1	MEFKCLIGHPEEEFVNLYRFRIGSKRKWPKMMDPSLSSSLKTCYKLNQTSRSFAAVQIA	60	1	MEFKCLIGHPEEEFVNLYRFRIGSKRKWPKMMDPSLSSSLKTCYKLNQTSRSFAAVQIA	60	1952	1952	374	374	1	374	MEFKCLIGHPEEEFVNLYRFRIGSKRKWPKMMDPSLSSSLKTCYKLNQTSRSFAAVQIA
2	VLSDDPFTCYHYAGLVGIGLSRLFSASBEFDPLVGEDTERANSMGLFLQKTNIRDLED	180	121	VLSDDPFTCYHYAGLVGIGLSRLFSASBEFDPLVGEDTERANSMGLFLQKTNIRDLED	180	1692	1692	416	416	10	416	VLSDDPFTCYHYAGLVGIGLSRLFSASBEFDPLVGEDTERANSMGLFLQKTNIRDLED
3	QSGGRFPQEVMSRYVKGFLPAKENPDILAVQCLBLITNALHHPDVITLRLRNQ	240	181	QSGGRFPQEVMSRYVKGFLPAKENPDILAVQCLBLITNALHHPDVITLRLRNQ	240	770.5	770.5	39.5	455	11	39.5	QSGGRFPQEVMSRYVKGFLPAKENPDILAVQCLBLITNALHHPDVITLRLRNQ
4	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	731	731	404	404	17	404	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
5	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	732	732	404	404	17	404	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
6	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	733	733	403	403	13	403	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
7	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	734	734	403	403	13	403	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
8	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	735	735	403	403	13	403	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
9	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	736	736	403	403	13	403	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
10	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	737	737	404	404	17	404	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
11	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	738	738	404	404	17	404	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
12	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	739	739	403	403	13	403	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
13	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	740	740	403	403	13	403	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
14	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	741	741	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
15	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	742	742	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
16	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	743	743	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
17	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	744	744	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
18	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	745	745	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
19	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	746	746	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
20	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	747	747	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
21	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	748	748	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
22	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	749	749	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
23	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	750	750	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
24	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	751	751	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
25	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	752	752	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
26	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	753	753	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME
27	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	241	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME	300	754	754	401	401	14	401	SVENFCATPQVMALATLACYNNQVKGAKKIRKGQAVLMDATNPAAKAIYOME

Sequence 170 , App
Sequence 221-233 , Sequence 3321 , Ap
Sequence 320955 , Sequence 44934 , Sequence 49024 , A
Sequence 203788 , Sequence 306328 , Sequence 43038 , Sequence 4986 , A
Sequence 52931 , A Sequence 133 , App
Sequence 28794 , Sequence 2 , Appli
Sequence 236999 , Sequence 18574 , A

Db 361 LSQVTEDVQVTGHE 374 ; LENGTH: 417
; TYPE: PRT
; ORGANISM: Human
; US-09-820-004-4

Query Match 98.4%; Score 1920.5; DB 3; Length 417;
Best Local Similarity 89.7%; Pred. No. 4.5e-184; Mismatches 0; Indels 43; Gaps 1;
Matches 374; Conservative 0; Mismatches 0; Indels 43; Gaps 1;

Oy 1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60
Db 1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60

Oy 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120
Db 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120

Oy 1 VLEDFT----- YCHYAGLVG 137
Db 1 VLEDFT----- YCHYAGLVG 137

Query Match 100.0%; Score 1952; DB 4; Length 374;
Best Local Similarity 100.0%; Pred. No. 2.6e-187; Mismatches 0; Indels 0; Gaps 0;
Matches 374; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60
1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60

Oy 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120
Db 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120

Oy 1 VLEDFT----- YCHYAGLVG 137
Db 1 VLEDFT----- YCHYAGLVG 137

Query Match 100.0%; Score 1952; DB 4; Length 374;
Best Local Similarity 100.0%; Pred. No. 2.6e-187; Mismatches 0; Indels 0; Gaps 0;
Matches 374; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60
1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60

Oy 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120
Db 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120

Oy 1 VLEDFT----- YCHYAGLVG 137
Db 1 VLEDFT----- YCHYAGLVG 137

Query Match 100.0%; Score 1952; DB 4; Length 374;
Best Local Similarity 100.0%; Pred. No. 2.6e-187; Mismatches 0; Indels 0; Gaps 0;
Matches 374; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60
1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60

Oy 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120
Db 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120

Oy 1 VLEDFT----- YCHYAGLVG 137
Db 1 VLEDFT----- YCHYAGLVG 137

RESULT 3 US-09-820-004-4

Sequence 4. Application US/09820004
; General Information:
; Patent No. US2001042418A1
; Sequence 5. Application US/09820004
; Patent No. US20020142418A1

Query Match 98.4%; Score 1920.5; DB 3; Length 417;
Best Local Similarity 89.7%; Pred. No. 4.5e-184; Mismatches 0; Indels 43; Gaps 1;
Matches 374; Conservative 0; Mismatches 0; Indels 43; Gaps 1;

Oy 1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60
Db 1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60

Oy 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120
Db 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120

Oy 1 VLEDFT----- YCHYAGLVG 137
Db 1 VLEDFT----- YCHYAGLVG 137

Query Match 98.4%; Score 1920.5; DB 3; Length 417;
Best Local Similarity 89.7%; Pred. No. 4.5e-184; Mismatches 0; Indels 43; Gaps 1;
Matches 374; Conservative 0; Mismatches 0; Indels 43; Gaps 1;

Oy 1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60
1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60

Oy 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120
Db 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120

Oy 1 VLEDFT----- YCHYAGLVG 137
Db 1 VLEDFT----- YCHYAGLVG 137

Query Match 98.4%; Score 1920.5; DB 3; Length 417;
Best Local Similarity 89.7%; Pred. No. 4.5e-184; Mismatches 0; Indels 43; Gaps 1;
Matches 374; Conservative 0; Mismatches 0; Indels 43; Gaps 1;

Oy 1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60
1 MEFVKCLIGHPEEFYLNVRFRIGGKRKVKMPKMDQDSLSSSLKTCYKYLNQTSRSFAVIOA 60

Oy 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120
Db 1 LDGEMRNACIVFYLNVRALDTEDDMTISVEKKVPLHNFHSFLYQDPWMFMESEKEKDRQ 120

Oy 1 VLEDFT----- YCHYAGLVG 137
Db 1 VLEDFT----- YCHYAGLVG 137